

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: July 19, 2002, 21:31:06 ; Search time 2528.22 Seconds
(without alignments)

18247.064 Million cell updates/sec

Title: US-09-836-410A-2
Perfect score: 3418
Sequence: 1 caagtaacacccgcaagatg.....atgcaataaattgttttggg 3418

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 13736207 seqs, 6748477542 residues

Total number of hits satisfying chosen parameters: 27472414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*
1: em_estba:*
2: em_esthum:*
3: em_estin:*
4: em_estnu:*
5: em_estov:*
6: em_estpl:*
7: em_estro:*
8: em_hic:*
9: gb_estl:*
10: gb_est2:*
11: gb_hic:*
12: gb_gss:*
13: em_gss_hum:*
14: em_gss_inv:*
15: em_gss_pln:*
16: em_gss_vrt:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	950.2	27.8	2758	11 AK005056	Mus musculus
2	754.4	22.1	1093	10 BM456794	AGENCOURT
3	714.2	20.9	1959	11 AK007755	Mus musculus
4	704.8	20.6	821	10 BM044197	BM044197 603621442
5	679.2	19.9	704	10 BG067031	BG067031 H3049G08-
6	673.4	19.7	689	9 BB616617	BB616617 BB616617
7	656.4	19.2	659	9 BB659255	BB659255 BB659255
8	649.8	19.0	675	9 AV270853	AV270853 AV270853
9	646.2	18.9	652	9 AW107262	AW107262 uml3c03.x
10	644.6	18.9	654	10 BF012472	BF012472 ux5903.y
11	629.6	18.4	673	9 BB577716	BB577716 BB577716
12	623.8	18.3	635	9 AW260482	AW260482 um80e10.x
13	623.4	18.2	625	10 BG080108	BG080108 H3049G08-
14	617.8	18.1	710	9 AI744486	AI744486 wF89n01.x
15	616.2	18.0	719	9 AU130763	AU130763 AU130763
16	610	17.8	710	9 BF163411	BF163411 60171776
17	608	17.8	617	9 BB478039	BB478039 BB478039

18	607.4	17.8	744	10	BG671648
19	602.2	17.6	945	10	BF179047
20	601	17.6	978	10	BM463359
21	591	17.3	657	10	BF472586
22	588.2	17.2	593	10	BG065941
23	580	17.0	600	10	BG805362
24	579	16.9	941	10	BG623888
25	575	16.8	603	10	BG079208
26	574.8	16.8	589	9	AW048763
27	573.4	16.8	626	10	BI662491
28	571.6	16.7	578	10	BI078250
29	561.4	16.4	629	10	BE300741
30	559.2	16.4	622	9	AW534169
31	559	16.4	625	10	BM236189
32	545.8	16.0	549	10	BE852629
33	545.2	16.0	684	9	AI177404
34	541.2	15.8	582	10	BM022195
35	539.6	15.8	615	10	BI966451
36	537	15.7	549	10	BF453530
37	535.2	15.7	931	10	BI687745
38	529.8	15.5	601	10	BF472327
39	528.8	15.5	666	10	D86662
40	518.8	15.2	572	11	AK017653
41	516.8	15.1	600	10	BI989826
42	507.8	14.9	519	10	BF450556
43	506.8	14.8	510	9	AA435048
44	505.4	14.8	826	10	BI150077
45	505.2	14.8	511	10	BE626970

ALIGNMENTS

RESULT 1
AK005056
LOCUS
DEFINITION
Mus musculus adult male liver cDNA, RIKEN full-length enriched library, clone:1300019C06;related to PUTATIVE N-TERMINAL ACETYLTRANSFERASE, full insert sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
AK005056
AK005056.1 GI:12836717
HTC; CAP trapper.
Mus musculus (strain:C57BL/6J) adult male liver cDNA to mRNA, clone:lib-RIKEN full-length enriched mouse cDNA library
clone:1300019C06.
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
1 (sites)
Carninci, P. and Hayashizaki, Y.
High-efficiency full-length cDNA cloning
Meth. Enzymol. 303, 19-44 (1999)
PUBMED
99279253
REFERENCE
2 (sites)
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes
Genome Res. 10 (10), 1617-1630 (2000)
JOURNAL
MEDLINE
20499374
PUBMED
11042159
REFERENCE
3 (sites)
Shibata, K., Itoh, M., Alzawa, K., Nagaoka, S., Sasaki, N., Carninci, P., Konno, H., Akiyama, J., Nishi, K., Kitsumai, T., Tashiro, H., Itoh, M., Sum, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A., Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K., Fujiwaki, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer
Genome Res. 10 (11), 1757-1771 (2000)
JOURNAL

AK005056 2758 bp mRNA linear HTC 19-JAN-2002
Mus musculus adult male liver cDNA, RIKEN full-length enriched library, clone:1300019C06;related to PUTATIVE N-TERMINAL ACETYLTRANSFERASE, full insert sequence.

AK005056
AK005056.1 GI:12836717
HTC; CAP trapper.
Mus musculus (strain:C57BL/6J) adult male liver cDNA to mRNA, clone:lib-RIKEN full-length enriched mouse cDNA library
clone:1300019C06.

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE
1 (sites)
Carninci, P. and Hayashizaki, Y.

High-efficiency full-length cDNA cloning

Meth. Enzymol. 303, 19-44 (1999)

PUBMED
99279253

REFERENCE
2 (sites)
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Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes

Genome Res. 10 (10), 1617-1630 (2000)

JOURNAL
MEDLINE
20499374

PUBMED
11042159

REFERENCE
3 (sites)
Shibata, K., Itoh, M., Alzawa, K., Nagaoka, S., Sasaki, N., Carninci, P., Konno, H., Akiyama, J., Nishi, K., Kitsumai, T., Tashiro, H., Itoh, M., Sum, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A., Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K., Fujiwaki, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer

Genome Res. 10 (11), 1757-1771 (2000)

JOURNAL

Qy 887 tgaagccagccctggacacagcagagatttatttaattcaattgtgcaaaatacat 946
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Qy 1067 tcaggcatacaaaagcaatgaacaaatttgtgaagcacttaagaaatgtcatgaattga 1126
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Qy 1127 gagacattttatagaataaccagatgaccagtttgactttcatatcatactgtatggaa 1186
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Qy 1307 tctgcagatgagaacaagaacacagcaggtgtatcacagcaacatgtctgacaagagct 1366
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Db 1839 GAAGAAATGCTTACACAGCAAGAGAGAGCTCAGAAAGGCTAAGGTAGAGAGAGAG 1898
Qy 1427 aaaaaatccgaagaaagcgcgaacggaactccgaaagaaagaaaggtatgatga 1486
Db 1899 AAGCACACAGAGGAGCGCAACAGAAACCAACCAAGAAAGAAAGAGAGAGAGA 1958
Qy 1487 gaagaatattgagggcccaagaagagcttaccctgagaactggccaaggttgaac 1546
Db 1959 AGAAGTTACCACTGGTCTAAGGAAGAACTTATTCCTGAAAAACCTAGAAAAGGTTGACAA 2018
Qy 1547 tccattggaagaagctattaaagtttttaacaccattgagaactgggtgaagaacaagt 1606
Db 2019 TCCATTAGACAGGCATCAAGTTCTCTCCTGCAAAACCCCTGCTGCTGAGAGCAT 2078
Qy 1607 agaaactcatcttttgcctttgagatctactttaggaagaaagaaagtttcttttgatgct 1666
Db 2079 TCATACCCACCTGCTGGCTTTGAAATATATTTTAGAAAGGGAAGTTTTCCTAATGTT 2138
Qy 1667 acaatcagtaagcgggatttgcctattgattcttagctatcctcctgctcagtgatgc 1726
Db 2139 GCAGTCTCTTAACAGAGCCTTTGCAATTTGAAGTAATATCTTGGTTACATGAATGCTT 2198
Qy 1727 gattgcactcttcttcttctgtgtgtgaagtaagagacttaccgaacacagttgaacagt 1786
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Qy 1787 attaaacaagaataatgaatcgtcttttggagcaacaaatccaaagaattttaaataaac 1846
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Qy 1847 ctcttgaagagggaattctgtattcattgcccacatagattatcagctgccaaaatgggtata 1906
Db 2319 TTTCTCTAGACACAAATGCTACTCTCTTCAGCATCTACTTCAGAGTGCTAAAATGATGTA 2378
Qy 1907 ttattagattcttcttagtcaaaacagcagcaatagagctggcgcaacacacttgatgac 1966
Db 2379 TTTCTGACAAAGTCAAGGCAAGAAAGCAATTTGCTACAGCCACTAGTTGGATGAAC 2438
Qy 1967 cctcaacaacagaacacttcagacttgcattggaagtggtggaagccttgtgtgtagtag 2026

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Qy 2087 ttatcttggcttttcagctcctcctggatacgaag 2121
Db 2559 ACTCACAGCTGCTTTCTCTGCTGCTCGCGGGAAG 2593
RESULT 2
BM456794 1093 bp mRNA linear EST 05-FEB-2002
LOCUS AGENCOURT_6404070 NIH_MGC_92 Homo sapiens cDNA clone IMAGE:5583707
5', mRNA sequence.
ACCESSION BM456794
VERSION BM456794.1 GI:18505834
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 1093)
AUTHORS NIH-MGC http://mgc.nci.nih.gov/.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: Life Technologies, Inc.
DNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM12347 row: d column: 12
High quality sequence stop: 695.
FEATURES
source
1..1093
Location/Qualifiers
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/db_xref="taxon:9606"
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/lab_host="DH10B (phage-resistant)"
/note="Organ: testis; Vector: pCMV-SPORT6; Site_1: NotI;
Site_2: SalI; Cloned unidirectionally; oligo-dT primed.
Average insert size 2.5 kb. Library enriched for
full-length clones and constructed by Life Technologies."
BASE COUNT 415 a 183 c 234 g 253 t
ORIGIN
Query Match 22.1%; Score 754.4; DB 10; Length 1093;
Best Local Similarity 91.5%; Pred. No. 5e-119;
Matches 820; Conservative 0; Mismatches 73; Indels 3; Gaps 2;
Qy 683 tggaaaggaggaaccccaaccacattactttgggtccagctactatttggcacagcattta 742
Db 1 TGGAAAGGAGGAACCAACCAACACATTTACTTTGGTCCAGTACTTGGCACAACATTA 60
Qy 743 tgataaattggtcagccatccattgtctctggaatacataataactgcaattgaaaglac 802
Db 61 TGACAAAATTGGTCAGCCATCTATTGCTTTGGAGTACATAAATACTGCTATTGAAGTAC 120
Qy 803 accaacaattgataagaactcttcttgaagaagctaaatctataagcatgctgggaatat 862
Db 121 ACCTACATTAATGAAGCTCTTTCTCTGGAAGCTAAAAATCTATAAGCATGCTGGAAATAT 180
Qy 863 taaagaagctgccagggtggatggatgaagccaggccctggacacagcagcagcagatttat 922

of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies).
Note: this is a NIH_MGC library.
BASE COUNT 310 a 145 c 168 g 198 t
ORIGIN

Query Match 20.6%; Score 704.8; DB 10; Length 821;
Best Local Similarity 92.8%; Pred. No. 1.5e-110; Indels 2; Gaps 2;
Matches 761; Conservative 0; Mismatches 57; Indels 2; Gaps 2;
QY 594 gataaagaagaggtggaactcgttagaagaactagttggttatgaaacttctctaaaa 653
Db 2 GACAAAGAAAGGTGCAATCATAGAGAGTGTAGTAGTTATGAAACCTCTCTAAAA 61
QY 654 agttgtgcctatttaaccccaatgatgtaggaagaggaagaaacctccaaccacattact 713
Db 62 AGCTGCGGTTATTAAACCCCAATGATGATGGAAGGAGGAACCAACCAACCACTACT 121
QY 714 tgggtccagtactattggcacagcattatgataaaattggtcagccattctgtctg 773
Db 122 TGGGTCCAGTACTTGGCACACATATTGACAAAATTGGTCAGCACTATGCTTTC 181
QY 774 gaatacataaactcgaattgaaagtacacacacattgatagaactcttcttctgtaaaa 833
Db 182 GAGTACATAAATCTGCTATTGAAAGTACACCTACATTAATGAACCTCTTCTCGTGAAA 241
QY 834 gctaaaaactataagcaatgctgggaatattaaagaagctccagggtggtatgaggaagcc 893
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QY 894 caggccctgacacagacagacagatttataatccaaagtgtgcaaaatacatgttaaaa 953
Db 302 CAGGCTTGGACACAGACAGACATTTTCACTCCAAATGTGCAAAATACATGCTAAAA 361
QY 954 gccaaactgtattaagagggctgaagaattgtgtccaaagtgttcacgggggaagaactcca 1013
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QY 1014 gcggtagagaacctgaatgaatcagtgatgtatgtgtccagacagagtggtctcaggca 1073
Db 422 CGGTAGAGAAATTTGAATGAATGCAAGTGTGTTCCAAACAGAAATGTGCCAGGCT 481
QY 1074 tacaagaactgaacaatttggtagacacttaagaatgtcatgaatttgagagacat 1133
Db 482 TATAAGCAATGATAAATTTGGTGAAGCACTTAAGAAATGTCATGAGATTGAGAGACAT 541
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QY 1254 ttcaagcagcgagaattgctattgagatcta-tttgaagcttcacacacccctctgac 1312
Db 662 TTCAAGCGCAAGAAATGCTATAGAGATCTATTTTGAAGCTTCATGACACCCCTTAC 721
QY 1313 agatgagaacaaagaacacagggctgtatagcaaaaacatgtctgacaaaagagctaaagaa 1372
Db 722 AGATGAGAATAAAGAACACCAAGCTGATACAGCAAAACATGTCTGACAAAGAGCTTAAAGAA 781
QY 1373 actcgtaataaaacaaagaag-agctcaaaagaagccca 1411
Db 782 CTAAGTATATAAACAAGAAAGAAAGCTCAAAAGAAAGCCCA 821

RESULT 5
BG067031/c
LOCUS
DEFINITION H3049G08-3 NTA Mouse 15K cDNA Clone Set Mus musculus linear EST 26-JAN-2001
H3049G08 3', mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

REFERENCE,
AUTHORS
TITLE
JOURNAL
COMMENT

FEATURES
source

BG067031
BG067031.1 GI:12549600
EST.
house mouse.
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 704)
Kargul,G.J., Dudekula,D.B., Qian,Y., Lim,M.K., Jaradat,S.A., Tanaka
,T.S., Carter,M.G. and Ko,M.S.H.
Verification and initial annotation of NIA mouse 15K cDNA clone set
Unpublished (2001)
Other_ESTS: H3049G08-5
Contact: George J. Kargul
Laboratory of Genetics
National Institute on Aging/National Institutes of Health
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
Email: cdna@lgsun.grc.nia.nih.gov
This clone set has been freely distributed to the community. Please
visit http://lgsun.grc.nia.nih.gov/cDNA/15K.html for details.
Plate: H3049 row: G column: 08
Seq primer: -21M13 Forward
High quality sequence stop: 704
POLYA=tes.

Location/Qualifiers
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/strain="C57BL/6J"
/db_xref="niaEST:H3049G08-3"
/db_xref="taxon:10090"
/clone="H3049G08"
/clone_lib="NIA Mouse 15K cDNA Clone Set"
/sex="Clones arrayed from a variety of cDNA libraries"
/dev_stage="Clones arrayed from a variety of cDNA libraries"
/lab_host="DH10B"
/note="Vector: pSPORT1; Site_1: SalI; Site_2: NotI; This clone is among a rearranged set of 15,247 clones from 11 embryo cDNA libraries (including preimplantation stage embryos from unfertilized egg to blastocyst, embryonic part of E7.5 embryos, extraembryonic part of E7.5 embryos, and E12.5 female mesonephros/gonad) and one newborn ovary cDNA library. Average insert size 1.5 kb. All source libraries are cloned unidirectionally with Oligo(dT)-Not primers. References include: (1) Genome-wide expression profiling of mid-gestation placenta and embryo using a 15,000 mouse developmental cDNA microarray, 2000, Proc. Natl. Acad. Sci. U S A, 97: 9127-9132; (2) Large-scale cDNA analysis reveals phased gene expression patterns during preimplantation mouse development, 2000, Development, 127: 1737-1749; (3) Genome-wide mapping of unselected transcripts from extraembryonic tissue of 7.5-day mouse embryos reveals enrichment in the t-complex and under-representation on the X chromosome, 1998, Hum Mol Genet 7: 1967-1978."
BASE COUNT 168 a 149 c 124 g 263 t
ORIGIN

Query Match 19.9%; Score 679.2; DB 10; Length 704;
Best Local Similarity 99.4%; Pred. No. 3.7e-106;
Matches 692; Conservative 0; Mismatches 3; Indels 1; Gaps 1;
QY 729 ttggcacagcattatgataaaattggtcagccatccattgtctctggaatacataaact 788
Db 695 TTTGCACAGCATTTATGATAAAATTTGGTCAGCCATCCATTGCTCTGGAATACATAAACT 636
QY 789 gcaattgaaagtacacacacattatgataaactcttctgtgaaagctaaataataag 848
Db 635 GCAATTGAAAGTACACCAACATTTGATGAACTCTTCTTGTAAAAGCTAAATCTAAG 576
QY 849 catgctgggaataataagaagctgccagggtggatgatgaagcccgccctggacaca 908
|||||

241	GTGAAAGTAAAGACTTACCCGAAACAGTTAGAACAGTATTAAACACAGAAATGAATCGTC	300
1810	tttttggagcaacaataccaagaattttaataaagaaacctttctgaaaaggaattctgatt	1869
301	TTTTTGGAGCAACAATCCAAAGAAATTTTAATGAACCTTTCTGAAAAGGAATTCGTATT	360
1870	catggccacatagattatcagctgcacaaatggtatattattattagattcttctagtcaca	1929
361	CATTGCCACATAGATTATCAGTGTCCAAATGGTATATTATTAGATTCTTCTAGTCAAA	420
1930	aacgagcaatagagctggcgacaaacacttgatgatccctcaccacagaacaccttcaga	1989
421	AACGAGCAATAGAGCTGGCGACAACTTCATGGATCCCTCACCACAGAAACCTTCAGA	480
1990	cttgcatggaagtgtggaagcctgtgtgtagctacgagactgttaagaagctg	2049
481	CTTGTCATGAGGTGTTCGAAGCCTTGTGTGATGGTGAAGCTAGGAGACTGTAAGAAGCTG	540
2050	ccgaagcctacagacaagtgtgtcataaagcttttcccttatgctgttgggttctatgcctc	2109
541	CCGAAGCCTACAGACGAAGTTGTCTAAGCTTTTCCCTTATGCTTTGGCTNTTCATGCCTN	600
2110	ctggtatcacgaagggatgaagatcacacagtgacacgagagatgcttcgagaacggaag	2169
601	CTGNATACGAAGAGGATATGAAGATCACAGTGAACGGAGATAGTCTGCAGAAACGGAG	660
2170	aactggccaatgaatctgaacatcatta	2198
661	AACTGGCNCATGAATCTGAACATCATTA	689
RESULT 7		
BB659255	659 bp mRNA linear	EST 26-OCT-2001
LOCUS	musculus cDNA full-length enriched, 13 days embryo heart Mus	
DEFINITION	musculus cDNA clone D330028A07 5', mRNA sequence.	
ACCESSION	BB659255.1	GI:16493078
VERSION	BB659255.1	
KEYWORDS	EST.	
SOURCE	house mouse.	
ORGANISM	Mus musculus	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
AUTHORS	Mammalia; Eutheria; Rodentia; Sclerognathi; Muridae; Murinae; Mus.	
	1 (bases 1 to 659)	
	Arakawa,T., Carninci,P., Fukuda,S., Furuno,M., Hanagaki,T., Hara,A.,	
	Hiramoto,K., Hori,F., Ishii,F., Ito,M., Kawai,J., Konno,H., Kouda	
	M., Koya,S., Matsuyama,T., Miyazaki,A., Nomura,K., Ohno,M., Sasaki	
	Okazaki,Y., Okido,T., Saito,R., Sakai,C., Sanjo,H., Suzuki,H.,	
	D., Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H.,	
	Tagami,M., Tagawa,A., Takahashi,F., Takeda,Y., Tanaka,T., Toyota,T.,	
	Muramatsu,M. and Hayashizaki,Y.	
	RIKEN Mouse ESTs (Arakawa,T., et al. 2001)	
TITLE	Unpublished (2001)	
JOURNAL	Contact: Yoshihide Hayashizaki	
COMMENT	Laboratory for Genome Exploration Research Group, RIKEN Genomic	
	Sciences Center(GSC), Yokohama Institute	
	The Institute of Physical and Chemical Research (RIKEN)	
	1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan	
	Tel: 81-45-503-9222	
	Fax: 81-45-503-9216	
	Email: genome-res@gsc.riken.go.jp,	
	URL:http://genome.gsc.riken.go.jp/	
	Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K., Itoh	
	M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.	
	Normalization and subtraction of cap-trapper-selected cDNAs to	
	prepare full-length cDNA libraries for rapid discovery of new	
	genes. Genome Res. 10 (10), 1617-1630 (2000)	
	wagi,K., Fujiwaka,S., Inoue,K., Togawa,Y., Izawa,M., Ohara,E.,	
	Watahiki,M., Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T., Matsura	
	S., Kawai,J., Okazaki,Y., Muramatsu,M., Inoue,Y., Kira,A. and	
	Hayashizaki,Y.	
	RIKEN integrated sequence analysis (RISA) system-384-format	
	Genome Res.	

0 (11), 1757-1771 (2000)	Konno,H., Fukunishi,Y., Shibata,K., Itoh,M., Carninci,P., Sugahara
Y. and Hayashizaki,Y.	Computer-based methods for the mouse full-length cDNA
encyclopedia: real-time sequence clustering for construction of a	nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
Kondo,S., Shinagawa,A., Saito,T., Kiyosawa,H., Yamanaka,I., Aizawa	K., Fukuda,S., Hara,A., Itoh,M., Kawai,J., Shibata,K. and
Hayashizaki,Y.	Computational Analysis of Full-Length Mouse cDNAs Compared with
Human Genome Sequences. Mamm. Genome. 12, 673-677 (2001)	Please visit our web site (http://genome.gsc.riken.go.jp) for
further details.	e mouse tissues.
FEATURES	Location/Qualifiers
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	prepared and sequenced in Mouse Genome Encyclopedia
	Project of Genome Exploration Research Group in Riken
	Genomic Sciences Center and Genome Science Laboratory in
	RIKEN. Division of Experimental Animal Research in Riken
	contributed to prepare mouse tissues. 1st strand cDNA was
	primed with a primer [5,
	GAGAGAGAGCGCGCAACTCGAGTTGTTTTTTTTTTTTT 3'], cDNA was
	prepared by using trehalose thermo-activated reverse
	transcriptase and subsequently enriched for full-length by
	cap-trapper. Second strand cDNA was prepared with the
	primer adapter of sequence [5,
	GAGAGAGAGATTCGAGTTGTAATTA


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Db 421 ACCAACAGAAACCTTCAGACTTGCATGGAGGTGTTGGAAGCCTTGTGTATGCTAGCCTA 480
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Db 481 GGAGACTGTAAAGAAGCTGCCGAAGCCTACAGAGCAAGTTGTCTATAAGCTTTTCCCTTAT 540
QY 2091 gcttgctttcatgctcctgctgatacagagagatgaagatcacagtggaacagat 2150
Db 541 GCTTTGGCTTTTCATGCTCCTCGATACGAGAGGATATGAAGATCACAGTGGAACGGAGAT 600
QY 2151 agttctgcagaacgaagaactgcccgaatgaaatctgaacatcattaaacaagcaaatg 2210
Db 601 AGTTCTGAGAACGGAAGACTGGCCATGAATCTGAACATCATTAACAAGCAAAATG 660
QY 2211 gaatgactttggacc 2225
Db 661 GAATGACTTTGGACC 675

RESULT 9
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LOCUS
DEFINITION
unl3c03.x1 Sugano mouse kidney mkia Mus musculus cDNA clone
IMAGE:2192164 3' similar to WP:Y50D7_164.A CE22298 ;, mRNA
sequence.
ACCESSION
AW107262
VERSION
AW107262.1 GI:6078062
KEYWORDS
EST.
SOURCE
house mouse.
ORGANISM
Mus musculus
REFERENCE
1 (bases 1 to 652)
AUTHORS
Marra,M., Hillier,L., Kucaba,T., Martin,J., Beck,C., Wylie,T.,
Underwood,K., Steptoe,M., Theising,B., Allen,M., Bowers,Y., Person
B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter
E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R.,
Waterston,R. and Wilson,R.
TITLE
The WashU-NCI Mouse EST Project 1999
JOURNAL
Unpublished (1999)
COMMENT
Contact: Marra M/WashU-NCI Mouse EST Project 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@wustl.edu
This clone is available royalty-free through LBNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: custom primer used
High quality sequence stop: 493.
MG1:1004616
Location/Qualifiers
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/clone_lib="Sugano mouse kidney mkia"
/sex="female"
/dev_stage="adult"
/lab_host="DH10B"
/notes="Organ: kidney; Vector: pME18S-FL3; Site_1: DraIII
(CACTGTGTG); Site_2: DraIII (CACATGTG); 1st strand cDNA
was primed with an oligo(dT) primer
[ANGTGGCCCTTTTCTTTTCTTTT]; double-stranded cDNA was
ligated to a DraIII adaptor [TGTGGCCTACTGG], digested
and cloned into distinct draIII sites of the pME18S-FL3
vector (5' site CACTGTGTG, 3' site CACATGTG). XhoI should
be used to isolate the cDNA insert. Size selection was
performed to exclude fragments <1.5kb. Library
constructed by Dr. Sumio Sugano (University of Tokyo
Institute of Medical Science). Custom primers for
sequencing: 5' end primer CTCTGCTCTAAAGAGCTCGG and 3' end
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BASE COUNT 155 a 138 c 114 g 244 t 1 others
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Best Local Similarity 99.4%; Pred. No. 1.6e-100;
Matches 648; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
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QY 825 ctgttaaaagcttaaaatctataagcatgctgggaatatttaaagaagctgccaggtgagtg 884
Db 592 CTGTGTAAGAGCTANAATCTATAAGCATGCTGGCAATATTAAAGAAAGCTGCCAGGTGGATG 533
QY 885 gatgaagcccgccctggagacacagacagacatttatttaattccaagtgtcaaaatac 944
Db 532 GATGAAGCCCGAGCCCTGGACACAGACAGACATTTATTAATTCGAAGTGTGCAAAATAC 473
QY 945 atgttaaaagcccaactgattaaagagggctgaagaaatgtgttccaagtttacgagggaa 1004
Db 472 ATGTTAAAAGCCCACTGATTAAGAGAGCTGAAGAAATGTGTTCGAAGTTTACGAGGGAA 413
QY 1005 ggaacttcagcggtagagaaacctgaatgaaatgcagtgatgtgtgttcagacagagtggt 1064
Db 412 GGAACCTTCAGCGGTAGAGAAACCTGAATGAATGCCAGTGTATGTGTTCAGACAGAGTGT 353
QY 1065 gctcaggcatcaaaagcaatgaacaaatgtgtgaagcacttaagaataatgtcatgaatt 1124
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QY 1125 gagagacattttatagaatacccgatgacccagtttgacccagtttgactttcactatcatgtatgag 1184
Db 292 GAGAGACATTTTATAGAAATCACCGATGACCGATGACCGATGACCGATGACCGATGACCG 233
QY 1185 aagatcaccccttagatcatatgtgagctattataaaactagaagatgacttcagacagcat 1244
Db 232 AGATCACCCCTTAGATCATATGTGGACTTATTAAAACTAGAAAGATGTACTTCGACAGCAT 173
QY 1245 ccattttactcaaacagcagcaaatgtctattgagatctatttgaagcttcactgacac 1304
Db 172 CCATTTTACTTCAAGCAGCGAGGATGCTATTGAGATCTATTGAGCTTTCATGACAAAC 113
QY 1305 cctctgacagatgagacaagaacacagcaggtgtgatacagcaaacatgtctgacaagag 1364
Db 112 CCTCTGACAAATGAGAAACAAAGAACACAGAGGCTGATACAGCAAAACATGCTGTGACAAAGAG 53
QY 1365 ctaagaagaactggttaataaaacaaagaagctcaaaagaagcccgagattg 1416
Db 52 CTAAGAAGAACTGCGTAAATAAAACAAAGAGAGCTCAAAAGAAAGACCCCATAG 1

RESULT 10
BF012472
LOCUS
DEFINITION
ux56g03.y1 Soares_NKWMd_mandible Mus musculus cDNA clone
IMAGE:3514324 5' similar to TR:Q9VWI2_Q9VWI2 CGI2202 PROTEIN. ;,
mRNA sequence.
ACCESSION
BF012472
VERSION
BF012472.1 GI:10712747
KEYWORDS
EST.
SOURCE
house mouse.
ORGANISM
Mus musculus
REFERENCE
1 (bases 1 to 654)
AUTHORS
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL
Unpublished (1997)
COMMENT
Contact: Robert Strausberg, Ph.D.
```


Email: cgapbs-re@mail.nih.gov
This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:1397204
Seq primer: -40RP from Gibco
High quality sequence stop: 471.
Location/Qualifiers
1. 654

FEATURES

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/note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site1: NotI; Site2: EcoRI; 1st strand cDNA was primed with a Not I - oligo(dT) primer (5' TGTTACCAATCTGAAGTGGAGCGGCCCTTAATTTTTTTTTTTT 3'), double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo."
BASE COUNT 228 a 118 c 137 g 168 t 3 others
ORIGIN

Query Match 18.9%; Score 644.6; DB 10; Length 654;
Best Local Similarity 98.9%; Pred. NO. 3e-100;
Matches 647; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

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QY 726 tatttggcacagcattatgataaaattggtcagccatccattgctctggaatacataaat 785
DB 61 TATTGGCACAGCATATGATAAAATTTGGTCAGCCATCATTTGCTCGAATACATAAAT 120

QY 786 actgcaattgaaagtacacccaacttgatagaactcttcttggtaaaagctaaatctat 845
DB 121 ACTGCAATTGAANGTACACCAACATTTGATAGAACTCTTTCTGTGTAAGCTAAATCTAT 180

QY 846 aagcatgctgggaattattaaagaagctgccaggtgtagtgatgaagccagccagccctggac 905
DB 181 AAGCATGCTGGGAATATTAAAGANGCTGAGGTGATGGATGAGTGAAGCCAGCCCTTGAC 240

QY 906 acagcagacagattatttaattccaagtgtgcacaaatacatgtttaaaagccaaacctgatt 965
DB 241 ACAGCAGACAGATTTATTATTTCAAGTGTGCATAAATACATGTTAAAGCCCAACCTGATT 300

QY 966 aaagaggtggaagaaatgttccaagtttacgaggggaagaaacttcagcggtagagaac 1025
DB 301 AAAGAGGCTGAAGAAATGTTTCCAAAGTTTACGAGGGAAGAACTTCAGCGGTAGAGAAC 360

QY 1026 ctgaatgaaatgcagtgatgtggttccagacagagtgctgcaggccatacaaaagcaatg 1085
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DB 421 AACAAATTTGTTGAAGCACTTTAAGAAATGTGATGAAATTTGAGAGACATTTATAGAAATC 480

QY 1146 accgatgaccagtttgactttcatcacactgtatgaggaagatcaccccttagatcatat 1205
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QY 1206 gtgacttattaaactagagaatgtactctgcagacagcatccattttacttcaaaagcagcg 1265
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RESULT 11
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LOCUS BB577716
DEFINITION BB577716 RIKEN full-length enriched, adult male medulla oblongata
ACCESSION BB577716
VERSION BB577716
KEYWORDS BB577716.2 GI:16449448
SOURCE EST.
ORGANISM house mouse.
MUS musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 673)
AUTHORS Arakawa,T., Carninci,P., Fukuda,S., Furuno,M., Hanagaki,T., Hara,A., Hiramoto,K., Hori,F., Ishii,Y., Ito,M., Kawai,J., Konno,H., Kouda,M., Koya,S., Matsuyama,T., Miyazaki,A., Nomura,K., Ohno,M., Okazaki,Y., Okido,T., Saito,R., Sakai,C., Sakai,K., Sano,H., Sasaki,D., Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagami,M., Tagawa,A., Takahashi,F., Takeda,Y., Tanaka,T., Toya,T., Muramatsu,M. and Hayashizaki,Y.
RIKEN Mouse ESTs (Arakawa,T., et al. 2001)
UNPUBLISHED (2001)
TITLE On Nov 30, 2000 this sequence version replaced gi:11474260.
JOURNAL Contact: Yoshihide Hayashizaki
COMMENT Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center(GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-Ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsr.riken.go.jp/
URL: http://genome.gsc.riken.go.jp/
Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K., Itoh,M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)
wagi,K., Fujiwaki,S., Inoue,K., Togawa,Y., Izawa,M., Ohara,E., Watahiki,M., Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T., Matsuura,S., Kawai,J., Okazaki,Y., Muramatsu,M., Inoue,Y., Kira,A. and Hayashizaki,Y.
RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)
Konno,H., Fukunishi,Y., Shibata,K., Itoh,M., Carninci,P., Sugahara,Y. and Hayashizaki,Y.
Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
Kondo,S., Shinagawa,A., Saito,T., Kiyosawa,H., Yamanaka,I., Alzawa,K., Fukuda,S., Hara,A., Itoh,M., Kawai,J., Shibata,K. and Hayashizaki,Y.
Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
Please visit our web site (http://genome.gsc.riken.go.jp/) for further details.
cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.
Location/Qualifiers
1. 673
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/sex="male"

FEATURES
source


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Db 515 GATGAAGCCAGCCCTGGACACAGACAGACAGATTATTAAATCCAGTGTGCAAAATAC 456
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Db 455 ATGTAAAGCCCAACCTGATTAAAGAGCGTGAAGAAATGTGTCCAAGTTTACAGGGAA 396
QY 1005 ggaactcagcgtgtagaagcctgaatgaatcgagtgatgtgttccagacagagtgt 1064
Db 395 GGAACCTTCTCGGTAGAGAACCTGAATGAATGCAGTGTGTGTGTTCCAGACAGAGTGT 336
QY 1065 gctcgggcatcaaaagcaatgaacaaatttggtgaagcacttaagaataatgcatagaatt 1124
Db 335 GCTCAGGCATACAAAGCAATGAACAAATTTGGTGAAGCACITTAAGAAATGTCATCAAAAT 276
QY 1125 gagagacattttatagaataaccagatgacagagtttgactttccatatactgtatgagg 1184
Db 275 GAGAGACATTTTATAGAATCACCAGTTCAGAGTTTGACTTTTCATACATACATGTTATGAGG 216
QY 1185 aagatcccttagatcatatgtgacttattaaactagaagatgtacttcgacagcat 1244
Db 215 AAGATCACCTTAGATCATATGTGACATTAAACCTAGAAGATGTACTTCGACAGCAT 156
QY 1245 caattttacttcaagcgcgagaattgctattgagatctattgaaagcttcacgacaac 1304
Db 155 CCATTTTACTTCAAGACGACGAGGATTGCTATTGAGATCTATTGAAGCTTCATGACAAAC 96
QY 1305 cctctgacagatgagaacaaagacagcagctgacacagaaacatgtctgcagaaagag 1364
Db 95 CCTCTGACAGATGAGACAAAGAACACACGAGGCTGATACAGCAAAATGTAAGACAAAGAG 36
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Db 35 CTAAGAAGCTGCGGAATAAACAAGAAGAGCTCA 1
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RESULT 13

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LOCUS
DEFINITION
H3049G08-5 NIA Mouse 15K cDNA Clone Set Mus musculus EST 26-JAN-2001
ACCESSION
BC080108
VERSION
BC080108.1 GI:12562676
KEYWORDS
EST.
SOURCE
Mus musculus
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 625)
Kargul, G.J., Dudekula, D.B., Qian, Y., Lim, M.K., Jaradat, S.A., Tanaka
T.S., Carter, M.G. and KO, M.S.H.
Verification and Initial annotation of NIA mouse 15K cDNA clone set
Unpublished (2001)
Other ESTs: H3049G08-3
Contact: George J. Kargul
Laboratory of Genetics
National Institute on Aging/National Institutes of Health
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
Email: cdna@igsun.grc.nia.nih.gov
This clone set has been freely distributed to the community. Please
visit http://igsun.grc.nia.nih.gov/cDNA/15k.html for details.
Plate: H3049 Row: G Column: 08
Seq primer: -21M13 Reverse
High quality sequence stop: 625
POLYA=NO.
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FEATURES

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Location/Qualifiers
1..625
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/dev_stage="Clones arrayed from a variety of cDNA
libraries"
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clone is among a rearrayed set of 15,247 clones from 11
embryo cDNA libraries (including preimplantation stage
embryos from unfertilized egg to blastocyst, embryonic
part of E7.5 embryos, extraembryonic part of E7.5 embryos
, and E12.5 female mesonephros/gonad) and one newborn
ovary cDNA library. Average insert size 1.5 kb. All
source libraries are cloned unidirectionally with Oligo(dT
)-Not primers. References include: (1) Genome-wide
expression profiling of mid-gestation placenta and embryo
using a 15,000 mouse developmental cDNA microarray, 2000,
Proc. Natl. Acad. Sci. U S A, 97: 9127-9132; (2)
Large-scale cDNA analysis reveals phased gene expression
patterns during preimplantation mouse development, 2000,
Development, 127: 1737-1749; (3) Genome-wide mapping of
unselected transcripts from extraembryonic tissue of
7.5-day mouse embryos reveals enrichment in the t-complex
and under-representation on the X chromosome, 1998, Hum
Mol Genet 7: 1967-1978."
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ORIGIN
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Db 1 AAAAATATATAGGAAGCCCTGGACTAAATACCCAGGGGACTCGTGCACAGAGGCTGCC 60

QY 482 cttaaaccttttctctggagagaagtttaaggaggtgttgtagaggttctctaagagtgaa 541
Db 61 CTTAAACTTTTATCTCGAGAGAGTTTANGAGAGTGTGTGGATAGGTTCCTAAGGATCAA 120

QY 542 ttccagcaaggggtgtccacctgtcttcaataacctgaggtctttatcacagagataaga 601
Db 121 TTTTCAGCAAGGGTGTCCACCTGTCTTCAATACCTTGAGGTCTTTATACAGAGATAAGA 180

QY 602 gaaggtggcaatcgtagaagaacttagttgattgaacacttctctaaaagttgtcg 661
Db 181 GAAGGTGGCAATCGTAGAAGAACTAGTAGTGTGTTATGAACACTTCTCTAAAAAGTTGTGC 240

QY 662 cctatttaaccccaatgatgatggaagaggaggaacccctcccaaccacattactttgggccc 721
Db 241 CCTATTTAACCCCAATGATGATGGAAGGAGGAACCTCCCAACCACTTACTTTGGGTCCA 300

QY 722 gtactattggccagcattatgatataaattggtcagccatccatctgtctggaatacat 781
Db 301 GTACTATTTGGCACACGATTTATGATAAAATTTGTCAGCCATCCATTTGCTCTGGAATACAT 360

QY 782 aaatactgcaattgaaagtacaccaacattgatagaactcttcttctgtaaaagctaaat 841
Db 361 AATACTGCAATTTGAAGATACACCAACATTTGATAGAACTCTTTCTTTGTAAGACTAAAT 420

QY 842 ctataagcatcgtgggaataattaaagaagctgccaggtggatggatgaagccagccct 901
Db 421 CTATAAGCATGCTGGGAATATTAAAGAAGCTGCCAGGTGGATGGAAGAGCCAGCCCT 480

QY 902 ggacacagacagacagatttataattccagagtgcaaaatacatgtttaaagccaaacct 961
Db 481 GGACACAGACAGACAGATTTTATTAATTTCCAAAGTGTGCAAAATACATGTTAAAGCCAACT 540

QY 962 gattaaagaggtggaagaaatgtgtcccaagtttacgaggggaaggaacttcagcggtaga 1021
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||||| GATTAAAGAGCTGAGAAATGCTTCCAGTTTACGAGGGAAGAACTTCAGCGGTAGA 600
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Qy 1022
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Db 601

RESULT 14
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sequence.
ACCESSION AI744486.1
VERSION AI744486.1
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 710)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-r@mail.nih.gov
Tissue Procurement: Elias Campo, M.D., Michael R. Emmert-Buck, M.D.,
Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.
cDNA Library Arraying: Greg Lannon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
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Seq primer: -40UP from Gibco
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was prepared from 12 pooled bulk tumor samples and primed
with a Not I - oligo(dT) primer. Double-stranded cDNA was
ligated to Eco RI adaptors (Pharmacia), digested with Not
I and cloned into the Not I and Eco RI sites of the
modified pT73 vector. Library went through one round of
normalization."
BASE COUNT 170 a 146 c 121 g 267 t 6 others
ORIGIN

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Matches 657; Conservative 0; Mismatches 53; Indels 1; Gaps 1;

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Db 650 GAGTACATAAATCTGCTATTGANAGTACA-CTACATTAATAGACNTCTTTCTCGTGANA 592
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Qy 834 gctaaaatctataagcatgctgggaattataaagaagctccaggtggatgaagacc 893

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VERSION AI130763.1
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ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 790)
AUTHORS Ota,T., Nishikawa,T., Suzuki,Y., Ishii,S., Saito,K., Kawai,Y.,
Yamamoto,J., Wakamatsu,A., Nakamura,Y., Nagai,T., Sugano,S. and
Isogai,T.
TITLE HRI human cDNA project
JOURNAL Unpublished (2000)
COMMENT Contact: Takao Isogai
Genomics Laboratory
Helix Research Institute
1532-3 Yana, Kisarazu, Chiba 292-0812, Japan
Tel: 81-438-52-3951
Fax: 81-438-52-3952
Email: genomics@hri.co.jp
HRI human cDNA project; 5'- & 3'-end one pass sequencing: Helix
Research Institute; cDNA library construction: Department of
Virology, Institute of Medical Science, University of Tokyo, and
Helix Research Institute.
FEATURES
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cells after 2-weeks retinoic acid (RA) induction"
BASE COUNT    250 a 136 c 155 g 246 t
ORIGIN

Query Match      18.0%; Score 616.2; DB 9; Length 790;
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Matches 718; Conservative 0; Mismatches 66; Indels 14; Gaps 5;

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QY 2062 gagcaagttgtcattgaagctttcccttctgtgtgtgtgtgtgtgtgtgtgtgtgtgt 2121
DB 181 GAGCAAAATTGTCTAAGCTTTTCCCTTATGCTTTGGCTTTCATGCTCCTCGATATGAAG 240

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DB 241 AGGATATGAAGATCACAGTTAATGGAGATAGTCTGCAGAAAGCTGAAGAACTGGCCAAATG 300

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us-09-836-410a-2.rst

Mon Jul 22 09:40:58 2002

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Date: Jul 20, 2002 3:07 AM

About: Results were produced by the GenCore software, version 4.5.
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Command line parameters:

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Database: EST:*

Database sequences: 13736207

Database length: -1841457050

Search time (sec): 1584.610000

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gb_htc:AK007755	+ 1374.00	2054.67	3.4e-105	1959	AK007755 Mus musculus 10 day c
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gb_est2:BE887347	+ 980.50	1490.50	8.9e-74	626	BE887347 601510175F1 NIH_MGC_7
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ACETYLTRANSFERASE, full insert sequence.
ACCESSION AK005056
VERSION AK005056.1 GI:12836717
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ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus;
1 (sites)
AUTHORS Carninci, P. and Hayashizaki, Y.
TITLE High efficiency full-length cDNA cloning
JOURNAL Meth. Enzymol. 303, 19-44 (1999)
MEDLINE 99279253
PUBMED 10349636
REFERENCE
2 (sites)
AUTHORS Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
TITLE Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new genes
JOURNAL Genome Res. 10 (10), 1617-1630 (2000)
MEDLINE 20499374
PUBMED 11042159
REFERENCE
3 (sites)
AUTHORS Shibata, K., Itoh, M., Aizawa, K., Nagao, S., Sasaki, N., Carninci, P.,
Konno, H., Akiyama, J., Nishi, K., Katsunai, T., Tashiro, H., Itoh, M.,
Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A.,
Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K.,
Fujikawa, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M.,
Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura, S., Kawai, J.,
Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
TITLE RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multichannel sequencer
JOURNAL Genome Res. 10 (11), 1757-1771 (2000)
MEDLINE 20530913
PUBMED 11076861
REFERENCE
4 (sites)
AUTHORS The RIKEN Genome Exploration Research Group Phase II Team and the
FANTOM Consortium.
TITLE Functional annotation of a full-length mouse cDNA collection
JOURNAL Nature 409, 685-690 (2001)
MEDLINE 20530913
PUBMED 11076861
REFERENCE
5 (bases 1 to 2758)
AUTHORS Adachi, J., Aizawa, K., Akahira, S., Akimura, T., Aono, H., Arai, A.,
Arakawa, T., Baldarelli, R., Bono, H., Brownstein, M., Bult, C.,
Carninci, P., Fukuda, S., Fukunishi, Y., Furuno, M., Hanagaki, T.,
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Saito, H., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D.,
Schriml, L., Shibata, K., Shibata, Y., Shinagawa, A., Shiraki, T.,
Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F.,
Tanaka, T., Tejima, Y., Toya, T., Yamamura, T., Yamanaka, I.,
Yasunishi, A., Yoshida, K., Yoshino, M., Muramatsu, M. and
Hayashizaki, Y.
```

```
Direct Submission
Submitted (10-JUL-2000) Yoshihide Hayashizaki, The Institute of
Physical and Chemical Research (RIKEN), Laboratory for Genome
Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
```

RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
Kanagawa 230-0045, Japan (E-mail: genome-res@cs.riken.go.jp,
URL: <http://genome.gsc.riken.go.jp/>, Tel: 81-45-503-9222,
Fax: 81-45-503-9216) shimizu@genome.gsc.riken.go.jp/ for

COMMENT

further details. cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. First strand cDNA was primed with a primer [5'-GAGACAGAGAGCGCGGCAACATCGAGTATTTTTTTTTTN 3'], cDNA was prepared by using thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. Second strand cDNA was prepared with the primer adapter of sequences [5'-GAGACAGAGAGAGTCCAGAGCTCAATTAATTAATTAACCCGCCCC 3']. cDNA was digested with SstI and SstI cloning sites. 5' end: SstI; 3' end:

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CDS

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polyA_signal

polyA_site

BASE COUNT	902 a	528 c	615 g	713 t
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alignment_scores:

alignment_scores:			
Quality:	2008.50	length:	591
Ratio:	3.915	Gaps:	1
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alignment_block:

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US-09-836-410A-1 x AK005056

Align seg 1/1 to: AK005056 from: 1 to: 2758

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      886  GAGAGCTTCAGCTTTATGAGGAGTGCAGCAAGCAGCAGCCCGAGAGCAGT 935

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Mon Jul 22 09:40:55 2002

JOURNAL
COMMENT

Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cqbbs-r@mail.nih.gov
Tissue Procurement: DCTD/DTP
CDNA Library Preparation: Ling Hong/Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone Distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>

Plate: LHCMI930 row: c column: 12
High quality sequence stop: 820.
Location/Qualifiers
1. 821

FEATURES

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/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5446979"
/clone_lib="NIH_MGC_40"
/tissue_type="Carcinoma, cell line"
/lab_host="DH10B (phage-resistant)"
/note="Organ: prostate; Vector: pOTB7; Site_1: XhoI;
Site_2: EcoRI; cDNA made by oligo-dT priming.
Directionally cloned into EcoRI/XhoI sites using the
following 5' adaptor: GGCACGAG(G). Library constructed by
Ling Hong in the laboratory of Gerald M. Rubin (University
of California, Berkeley) using ZAP-cDNA synthesis kit
(Stratagene) and Superscript II RT (Life Technologies).
Note: this is a NIH_MGC Library."
BASE COUNT 310 a 145 c 168 g 198 t
ORIGIN

alignment_scores:
Quality: 1401.00 Length: 274
Ratio: 5.151 Gaps: 2
Percent Similarity: 99.270 Percent Identity: 98.905
alignment_block:
US-09-836-410A-1 x BM044197
Align seg 1/1 to: BM044197 from: 1 to: 821
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2 CACAAAGAAAGGTTGGCAATCATAGAGAGTTAGTAGTTTATGAAC 51
79 rSerLeuLysSerCysArgLeuPheAsnProAsnAspGlyLysGluG 96
52 CTCCTAAAGAGCTCCGGTTATTATTAACCCCAATGATGGAAGAGG 101
96 LuProProThrThrLeuLeuTrpValGlnTyrTyrLeuAlaGlnHisTyr 112
102 AACCCACCACCATCTTTTGGGTCAGTACTTGGCACACATTTAT 151
113 AspLysIleGlyGlnProSerIleAlaLeuGluTyrIleAsnThrAla 129
152 CACAAATTTGGTCAGCCATCTATTCTTTGGAGTACATAAATACTG 201
129 eGluSerThrProThrLeuIleGluLeuPheLeuValLysAlaLys 146
202 TGAAGATACACCTACATTAATAGAACTCTTCTCGTGAAGCTAAAT 251
146 YrLysHisAlaGlyAsnIleLysGluAlaAlaArgTrpMetAspGlu 162
252 ATACGATGCTGGAATATTAAAGAGCTGCAAGGTGGATGGATGAG 301
163 GlnAlaLeuAspThrAlaAspArgPheIleAsnSerLysCysAlaL 179
302 CAGGCTTGGACACAGCAGACAGATTTATCAACTCCAAATGTGCA 351
179 rMetLeuLysAlaAsnLeuIleLysGluAlaGluMetCysSerLys 196
352 CATGCTAAAGAGCACTGATTAAGAGAGCTGAAGAAATGTGCTCAA 401

193 CysSerLysPheThrArgGluGlyThrSerAlaValGluAsnLeuAnG 209
302 TGCTCAAGATTTACAAGGAAGAACATCAGCGGTAGAGAAATTTGA 351
209 uMetGlnCysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLys 226
352 AATGCATGATGTGGTTCCAAACAGAAATGTGCCAGGCTTATTAAG 401
226 eAsnLysPheGlyGluAlaLeuLysCysHisGluIleGluArgHis 242
402 TGAATAAATTTGGTGAACACTTAAGAAATGTCATGAGATTGAGAC 451
243 PheIleGluThrAspAspGlnPheAspPheHisThrTyrCysMet 259
452 TTTATGAATACCTGATGATGACCCAGTTTACCTTTCATACATAC 501
259 gLysIleThrLeuArgSerTyrValAspLeuLeuLysLeuGluAsp 276
502 GAAGATTACCTTATGATCATATGTGGACTTTATTAAGACTAGAGAT 551
276 euArgGlnHisProPheThrPheLysAlaAlaArgIleAlaIleG 292
552 TTCGACGACATCCATTTTACTTCAAGCAGCAGCAAGATTTGCTAT 601
293 TyrLeuLysLeuHisAspAsnProLeuThrAspGluAsnLysGlu 309
602 TATTGAAGCTTTCATGACACCCCTTACAGATGAGAAATAAGACAG 651
309 uAlaAspThrAlaAsnMetSerAspLysGluLeuLysLysLeuArg 326
652 AGCTGATACAGCAACATCTCTGACAAAGAGCTAAAGAGCTACGTA 701
326 ysGlnArgArgAlaGlnLysLysAlaGlnIleGluGluGluLys 342
702 KACAAAGAGAGCTCAAAAGAAAGCCAGATAGAAAGAGAGAAAAA 751
343 AlaGluLysGluLysProGlnArgAsnProLysLysLysLysAsp 359
752 GCCGAAAGAAAGAACCCAGCAGAGAAATCAGAAAGAGAGAGGAT 801
359 pAspGluGluIleGlyGlyProLysGluGluLeuIleProGluLys 375
802 TGATGANGAGATAGGAGGCTCAAAAGAAAGAACTATTTCAGAGA 851
376 AlaLysValGluThrProLeuGluGluAlaIleLysPheLeuThr 391
852 GCCAAGGNTGAAAGCTCCATTTGGGAGAGAGCTTATTAATTTT 901
391 oLeuLysAsn.....LeuValLysAsnLysIleGluThrHisLeu 405
902 GGTGAAGAAAGCTGGGTCAAGCAACAGAAATAGAGGACCTCATCT 951
406 Ala...PheGlu...IleTyrPheArgLysGluLysPheLeuMet 419
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DEFINITION 603621442F1 NIH_MGC_40 Homo sapiens cDNA clone IMAGE:5446979 5',
mRNA sequence.
ACCESSION BM044197
VERSION BM044197.1 GI:16773464
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 821)
AUTHORS NIH-MGC <http://mgi.nci.nih.gov/>.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

Mon Jul 22 09:40:55 2002

alignment_block:

US-09-836-410A-1 x AK007755

Align seg 1/1 to: AK007755 from: 1 to: 1959

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912  GAGAGGCTTCAGCTTTATGAGGAGTGCAGAACGAGCAGCCGCCAGAGAGT  961
19  uValProArgLysLeuProLeuAsnPheLeuSerGlyGluLysPheLysG  36
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
962  TTCACCCAGGAGGCTGCGCTGAGTTTCGCCCCAGGTAAGAAGTTTCGAG  1011
36  LuCysLeuAspArgPheLeuArgMetAsnPheSerLysGlyCysProPro  52
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1012  AACTCATGGATAAGTTCCTGAGACCAACTTTAGCAAAAGGCTGTCCACCT  1061
53  ValPheAsnThrLeuArgSerLeuTyrArgAspLysGluLysValAlaI  69
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1062  CTGTTCTACTACTTTGAATACCTTGATGATGATGATGATGATGATGAT  1111
69  eValGluGluLeuValValGlyTyrGluThrSerLeuLysSerCysArgL  86
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1112  AATCCAGGAACCTGTTACTAATTATGAAGCCTCTCTTAAATGAATGGCT  1161
86  euPheAsnProAsnAspAspGlyLysGluGluProThrThrLeuLeu  102
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1162  APTTTTAGCCCTTATGAGAACGGGAGAGAACCCCAACCACTCTAATC  1211
103  TrpValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyGlnProSe  119
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1212  TGGTTCAGTATTCTTCCGACAGCATTTATGATAAATTTGGCAGTATT  1261
119  rIleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrProThrLeuI  136
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1262  TCTGGCTTTGGAATATATTAATGCTGTAATTGCTAGTACTCAACTTAA  1311
136  leGluLeuPheLeuValLysAlaLysIleTyrLysHisAlaGlyAsnIle  152
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1312  TAGAACTATTCTACATGAAAGCAAAANTTACAGCATATGGGTAACTC  1361
153  LysGluAlaAlaArgTrpMetAspGluAlaGlnAlaLeuAspThrAlaAs  169
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1362  AAAGAGCCGCACAGTGGATGATGATGAGGCACAGTCTTTGGACAGCGCTGA  1411
169  pArgPheIleAsnSerLysCysAlaLysTyrMetLeuLysAlaAsnLeuI  186
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1412  CAGGTTTCATCAATTCCAATGTGCCAAATACATGCTTCGAGCAATATGA  1461
186  leLysGluAlaGluMetCysSerLysPheThrArgGluGlyThrSer  202
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1462  TAAAGACAGCAGAGAAATGTGCTCCAGGTTCCAAAGGCAAGCAATCT  1511
203  AlaValGluAsnLeuAsnGluMetGlnCysMetTrpPheGlnThrGluCy  219
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1512  GCCATGGAGAACTGAATGAAATGCACTGTATGTGTTTGACAGCGAGTG  1561
219  sAlaGlnAlaTyrLysAlaMetAsnLysPheGlyGluAlaLeuLysLysC  236
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1562  CATTTCTGCCTATACGCGCTGGGAGATATGGGATGCTTGCTTGAAGAGT  1611
236  yHisGluIleGluArgHisPheIleGluIleThrAspAspGlnPheAsp  252
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1612  GCCATGAAGTAGAGAGCACTTTCTTGAGATAACCGATGATCATGTTGAC  1661
253  PheHisThrTyrCysMetArgLysIleThrLeuArgSerTyrValAspLe  269
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1662  TTCCATACATCTACATGAGAAAGATGACCCCTCCGGTCTTATGTTGCCT  1711
269  uLeuLysLeuGluAspValLeuArgGlnHisProPheTyrPheLysAlaA  286
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1712  CTTGAGATTAGAAGATCTCTCAGAGACATACATTTTATTTTCAAGGCTG  1761
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1762  CTGATCAGCAATTTGAATATATTTGAATTTACATGATAACCTTTAAACC  1811
303  AspGluAsnLysGluHisGluAlaAspThrAlaAsnMetSerAspLysG  319
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1812  ATGACAGCAAAACACAGACATAGATTTCAGAAAACCTGTACAGCCAAAGA  1861
319  uLeuLysLysLeuArgAsnLysGlnArgArgAlaGlnLysLysAlaGlnI  336
|||||  |||||||  |||||||  |||||||  |||||||  |||||||  |||||||
1862  AATCAAGAAAATGCTTAGCAAGCAAGAGAGCTCAGAAAAAGGCTAAGG  1911
336  leGluGluLysLysAsnAlaGluLysGluLysProGlnArgAsn  351
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1912  TAGAAGAAGAGAGAAAGCACACAGAAAGCGACGCCCAACAGAAAAAAC  1958
seq_name: gb_est2.BG067031

seq_documentation_block:
LOCUS      BG067031
DEFINITION H3049G08-3 NIA Mouse 15K cDNA Clone Set Mus musculus cDNA clone
ACCESSION  BG067031
VERSION    BG067031.1
KEYWORDS   EST.
SOURCE     house mouse.
ORGANISM   Mus musculus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE  1 (bases 1 to 704)
AUTHORS   Kargul, G.J., Dudekula, D.B., Qian, Y., Lim, M.K., Jaradat, S.A., Tanaka
            , T.S., Carter, M.G. and Ko, M.S.H.
TITLE     Verification and initial annotation of NIA mouse 15K cDNA clone set
JOURNAL   Unpublished (2001)
COMMENT   Other ESTs: H3049G08-5
            Contact: George J. Kargul
            Laboratory of Genetics
            National Institute on Aging/National Institutes of Health
            333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
            Email: cdna@lgsun.grc.nia.nih.gov
            This clone set has been freely distributed to the community. Please
            visit http://lgsun.grc.nia.nih.gov/cDNA/15k.html for details.
            Plate: H3049 row: G column: 08
            Seq primer: -21M13 Forward
            High quality sequence stop: 704
            POLYA=Yes.

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         /lab_host="DH10B"
         /note="Vector: pSPORT1; Site.1: SalI; Site.2: NotI; This
         clone is among a rearrayed set of 15,247 clones from 11
         embryo cDNA libraries (including preimplantation stage
         embryos from unfertilized egg to blastocyst, embryonic
         part of E7.5 embryos, extraembryonic part of E7.5 embryos
         , and E12.5 female mesonephros/gonad) and one newborn
         ovary cDNA library. Average insert size 1.5 kb. All
         source libraries are cloned unidirectionally with Oligo(dT
         )-Not primers. References include: (1) Genome-wide
         expression profiling of mid-gestation placenta and embryo
         using a 15,000 mouse developmental cDNA microarray, 2000,
         Proc. Natl. Acad. Sci. U S A, 97: 9127-9132; (2)
         Large-scale cDNA analysis reveals phased gene expression
         patterns during preimplantation mouse development, 2000,
         Development, 127: 1737-1749; (3) Genome-wide mapping of
```

unselected transcripts from extraembryonic tissue of 7.5-day mouse embryos reveals enrichment in the t-complex and under-representation on the X chromosome, 1998, Hum Mol Genet 7: 1967-1976."

BASE COUNT 168 a 149 c 124 g 263 t
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Quality: 1180.00 Length: 231
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Percent Similarity: 99.567 Percent Identity: 99.567

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US-09-836-410A-1 x BG067031/rev ..

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692 GCACAGCATTATGATAAAATTGGTCAGCCATCCATTCTCTGGAATACAT 643
125 eAsnThrAlaIleGluSerThrProThrLeuIleGluLeuPheLeuVal 142
|||||
642 AAATCTGCATTTGAAAGTACACCAACATTGATAGAACTCTTCTTGTA 593
142 ysAlaLysIleTyrLysHisAlaGlyAsnIleLysGluAlaAlaArgT 158
|||||
592 AAGCTAAATCTATAAGCATGCTGGGAATATTAAAGAGCTGCCAGGTGG 543
159 MetAspGluAlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSer 175
|||||
542 ATGGATGAAGCCAGGCCCTGGACAGCAGACAGATTTTATTATTCCAA 493
175 sCysAlaLysTyrMetLeuLysAlaAsnLeuIleLysGluAlaGluGlu 192
|||||
492 GTGTGCAAAATACATGTTTAAACCCCAACCTGATTAAAGAGCTGAAGAA 443
192 etCysSerLysPheThrArgGluGlyThrSerAlaValGluAsnLeuAsn 208
|||||
442 TGTGTTCCAAAGTTACGAGGGAAGAACTTCACGCGTAGAGAACCTGAT 393
209 GluMetGlnCysMetThrPheGlnThrGluCysAlaGlnAlaTyrLysAl 225
|||||
392 GAATTCAGTGTATGTGTTCCAGACAGAGTGTGCTCAGGCATACAAAGC 343
225 aMetAsnLysPheGlyGluAlaLeuLysLysCysHisGluIleGluArgH 242
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342 AATGAACAAATTTGGTGAAGCACTTAAGAAATGTCATGAATTTGAGAC 293
242 isPheIleGluIleThrAspAspGlnPheAspPheHisThrTyrCysMet 258
|||||
292 ATTTTATGAAATACCCGATGACCGATTTTGACTTTCATACATCTGTATG 243
259 ArgLysIleThrLeuArgSerTyrValAspLeuLysLeuGluAspVa 275
|||||
242 AGGAAGTACCCTTAGATCATATGTGGACTTATTAAAACTAGAAGATGT 193
275 leuArgGlnHisProPheTyrPheLysAlaAlaArgIleAlaIleGluI 292
|||||
192 ACTTCACAGCATCCATTTTACTTCAAGCAGCAGGAGTGTGTTTATGAGA 143
292 leTyrLeuLysLeuHisAspAsnProLeuThrAspGluAsnLysGluHis 308
|||||
142 TCTATTGAAAGCTTCATGACAAACCTCTGACAGATGAGAACAAAGAAC 93
309 GluAlaAspThrAlaAsnMetSerAspLysGluLeuLysLysLeuArgAs 325
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92 GAGGTGATACAGCAACATGTCTCACAAGAGAGCTTAAAGAAATG.CGTAA 44
325 nLysGlnArgArgAlaGlnLysLysAlaGlnIleGluGlu 339
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43 TAAACAAAGAGAGCTCAAAAGAAAGCCAGCATAGAGAAAGAG 1

seq_name: gb_est1:AI744486

seq_documentation_block:

LOCUS AI744486

DEFINITION wf89h01.x1 NCI_CGAP_Co3 Homo sapiens cDNA clone IMAGE:2362801 3'

similar to TR:074985 074985 N-TERMINAL ACETYLTRANSFERASE 1. ; mRNA

sequence.

ACCESSION: AI744486

VERSION: AI744486

KEYWORDS: EST.

SOURCE: AI744486.1 GI:5112774

ORGANISM: human.

REFERENCE: Homo sapiens

Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

1 (bases 1 to 710)

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

Tissue Procurement: Elias Campo, M.D., Michael R. Emmert-Buck, M.D.

, Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D.

cDNA Library Arraying: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 854 Std Error: 0.00

Seq Primer: -400P from Glibco

High quality sequence stop: 462.

FEATURES

Location/Qualifiers

1..710

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/db_xref="taxon:9606"

/clone="IMAGE:2362801"

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/sex="pooled"

/tissue_type="colon"

/lab_host="DH10B"

/note="Vector: pT73D-Pac (Pharmacia) with a modified

polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA

was prepared from 12 pooled bulk tumor samples and primed

with a Not I - oligo(dT) primer. Double-stranded cDNA was

ligated to Eco RI adaptors (Pharmacia), digested with Not

I and cloned into the Not I and Eco RI sites of the

modified pT73 vector. Library went through one round of

normalization."

BASE COUNT 170 a 146 c 121 g 267 t 6 others

ORIGIN

alignment_scores:
Quality: 1176.00 Length: 237
Ratio: 5.091 Gaps: 0
Percent Similarity: 97.468 Percent Identity: 96.203
alignment_block:
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103 TrpValGlnTyrLeuAlaGlnHisTyrAspLysIleGlyGlnProse 119
|||||
710 TGGGTCAAGTACTTGGCACACCATATGACAAATTTGTCAGCCATC 661
119 rleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrProThrLeuI 136
|||||
660 TANTGCTNTGGAGTACATAATACTGCTATTGANAGTACACT.ACATTAA 612
136 leGluLeuPheValLysAlaLysIleTyrLysHisAlaGlyAsnIle 152

DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the T.M.A.G.E. Consortium/LLNL at:

http://image.llnl.gov
plate: LLCM1637 row: p column: 10
High quality sequence stop: 724.

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59. The sixtieth column is labeled "SOURCE".	
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86. The eighty-seventh column is labeled "FEATURES".	
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95. The ninety-sixth column is labeled "SOURCE".	
96. The ninety-seventh column is labeled "FEATURES".	
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98. The ninety-ninth column is labeled "FEATURES".	
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/note="Organ: placenta; Vector: pNMR-L
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5' and 3' adaptors were used in cloning
adaptor sequence: 5'-CAGCGCATTTATGGCC
sequence: 5'-ATTCTAGAGCGAGGCGGCGAC
(where B = A, C, G and N = A, C, G
insert size 1.3 kb (range 0.5-4.0 kb)).
contained inserts by PCR. This library
full-length clones and was constructed
Laboratories (Palo Alto, CA). Note: the
Library."

Library.	358 a	172 c	211 g	200 t
BASE COUNT				
ORIGIN				

alignment_scores:		
Quality:	1170.50	Length: 286
Ratio:	4.608	Gaps: 5
Percent Similarity:	88.811	Percent Identity: 82.168

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PS-09-836-410A-1 x BG623888

Align seg 1/1 to: BG623888 from: 1 to: 941

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45  AGGAACCAACCAACACATTACTTTGGTGCCAGTACTACTTGGCAACA 94
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
111  sTyAspLysIleGlyGlnProSerIleAlaLeuGluTyrIleAsnThrA 128
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
95  TTATGACAAAATTGGTCAGCCATCTATTGCTTGGAGTACATAAATCTG 144
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
128  laIleGluSerThrProThrLeuIleGluLeuPheLeuValLysAlaLys 144
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
145  CTATTGAAGTACACCTTACATTAATAGAACTCTTCTCGTGAAGACTAAA 194
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
145  IleTyrLysHisAlaGlyAsnIleLysGluAlaAlaArgTrpMetAspGl 161
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
195  ATCTATAAGCATGCTGSAATAATTAAAGAAGCTGCAAGGTGGATGATGA 244
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
161  uAlaGlnAlaLeuAspThrAlaaspArgPheIleAsnSerLysCysAlaL 178
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
245  GGCCAGCGCCTTGACACACGACACAGATTATCAACTCCAAATGTGCCAA 294
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178  ySTyrMetLeuLysAlaAsnLeuIleLysGluAlaGluMetCysSer 194
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
295  AATACATGCTAAAAGCCAACTGATTAAAGACGCTGAAGAAATGTGCTCA 344
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
195  LysPheThrArgGluGlyThrSerAlaValGluAsnLeuAsnGluMetGl 211
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
345  AAGTTTACAAGGGAAGCAACATCATCGCGTGAAGAAATTTGAATGAAATGCA 394
    : : : : : : : : : : : : : : : : : : : : : : : : : : : :
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 DEFINITION mRNA sequence.

ACCESSION BG623888
VERSION BG623888.1 GI:13675259
KEYWORDS EST.
human

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human.	Homo sapiens	
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	

REFERENCE
1 (bases 1 to 941)
Mammalian, Domestic
NIH-MGC <http://mgc.nci.nih.gov/>.
AUTHORS
National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE
Unpublished (1999)
JOURNAL

CONTACT: Robert Strausberg, Ph.D.
Contact: Robert Strausberg, Ph.D.
Email: cgaaps@mail.nih.gov
Tissue Procurement: CLONTECH Laboratories, Inc.
CDNA Library Preparation: CLONTECH Laboratories, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (I

ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 625)
 REFERENCE Kargul, G.J., Dudekula, D.B., Qian, Y., Lim, M.K., Jaradat, S.A., Tanaka
 AUTHORS T.S., Carter, M.G. and Ko, M.S.H.
 TITLE Verification and initial annotation of NIA mouse 15K cDNA clone set
 JOURNAL Unpublished (2001)
 COMMENT Other_ESTS: H3049G08-3
 Contact: George J. Kargul
 Laboratory of Genetics
 National Institute on Aging/National Institutes of Health
 333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
 Email: cdna@igsun.grc.nia.nih.gov
 This clone set has been freely distributed to the community. Please
 visit <http://igsun.grc.nia.nih.gov/cDNA/15k.html> for details.
 Plate: H3049 row: G column: 08
 Seq primer: -21M13 Reverse
 High quality sequence stop: 625
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 ORIGIN

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 US-09-836-410A-1 x BG080108 ..
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 22 qLysLeuProLeuAsnPheLeuSerGlyGluLysPheLysGluCysLeuA 39
 52 AAGGCTGCCCTTAACTTTTATCTCGAGAGAGATTAAAGGAGTGTGTGG 101

39 spArgPheLeuArgMetAsnPheSerLysGlyCysProProValPheAsn 55
 102 ATAGGTTCTTAAAGTGAATTTTCAGCAAGGGCTGTCCACCTGTCTTCAAT 151
 56 ThrLeuArgSerLeuTyrArgAspLysGluLysValAlaIleValGluGl 72
 152 ACCTTGAGGTCCTTATACAGAGATAAGAGAGAGTGGCAATTCGTAGAGA 201
 72 uLeuValValGlyTyrCluThrSerLeuLysSerCysArgLeuPheAsnP 89
 202 ACTAGTAGTTGGTTATGAAACTTCTCTAAAGTGTGCGCTATTAAACC 251
 89 roAsnAspAspGlyLysGluGluProProThrThrLeuLeuTrpValGln 105
 252 CCAATGATGATGGAAGAGAGAGAGCTCCACCAACATTTACTTTGGGTCCAG 301
 106 TyrTyrLeuAlaGlnHisTyrAspLysIleGlyGlnProSerIleAlaLe 122
 302 TACTATTGGCACAGCAATATGATAAAATTTGGTCAGCCATCCATGCTCT 351
 122 uGluTyrIleAsnThrAlaIleGluSerThrProThrLeuIleGluLeuP 139
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 139 heLeuValLysAlaLysIleTyrLysHisAlaGlyAsnIleLysGluAla 155
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 156 AlaArgTrpMetAspGluAlaGlnAlaLeuAspThrAlaAspArgPheIl 172
 452 GCCAGGTGGATGGATGAAGCCAGCCCTGGACACAGACAGACATTTAT 501
 172 eAsnSerLysCysAlaLysTyrMetLeuLysAlaAsnLeuIleLysGluA 189
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 189 iaGluGluMetCysSerLysPheThrArgGluGlyThrSerAlaValGlu 205
 552 CTGAAGAAATGTGTTCGAAGTTTACGAGGGAAGAACTTCAGCGGTAGAG 601
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 seq_documentation_block:
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 sequence.
 ACCESSION AW260482
 VERSION AW260482.1 GI:66333463
 KEYWORDS EST.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 635)
 REFERENCE Marra M/WashU-NCI Mouse EST Project 1999
 AUTHORS Contact: Marra M/WashU-NCI Mouse EST Project 1999
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: mouseest@watson.wustl.edu
 TITLE The WashU-NCI Mouse EST Project 1999
 JOURNAL Unpublished (1999)
 COMMENT Other_ESTS: um80e10.y1

REFERENCE:
AUTHORS.

1 (bases 1 to 988)
 Aharawo, T., Carninci, P., Fukuda, S., Furuno, M., Hanagaki, T., Hara, A.,
 Hiramoto, K., Hori, F., Ishii, Y., Ito, M., Kawai, J., Konno, H., Kouda
 K., Koya, S., Matsuyama, T., Miyazaki, A., Nomura, K., Ohno, M.,
 Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki
 D., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H.,
 Tagami, M., Tagawa, A., Takahashi, F., Takeda, Y., Tanaka, T., Toyota, T.,
 Muramatsu, M., and Hayashizaki, Y.
 RIKEN Mouse ESTs (Arakawa, T., et al. 2001)
 Unpublished (2001)
 On Nov 1, 1999 this sequence version replaced gi:6179369.
 Contact: Yoshihide Hayashizaki
 Laboratory for Genome Exploration Research Group, RIKEN Genomic
 Sciences Center(GSC), Yokohama Institute
 The Institute of Physical and Chemical Research (RIKEN)
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
 Tel.: 81-45-503-9222

TITLE	JOURNAL	COMMENT
1. The Role of the Teacher in the Classroom	Journal of Educational Research	1980, Vol. 83, No. 1, pp. 1-10
2. The Impact of Technology on Education	Journal of Educational Technology	1985, Vol. 10, No. 2, pp. 1-15
3. The Importance of Parental Involvement	Journal of Educational Psychology	1990, Vol. 82, No. 3, pp. 1-12
4. The Effect of Teacher Expectations on Student Achievement	Journal of Educational Research	1995, Vol. 98, No. 4, pp. 1-18
5. The Role of the School in the Community	Journal of Educational Research	2000, Vol. 103, No. 5, pp. 1-20
6. The Impact of Teacher Education on Student Achievement	Journal of Educational Research	2005, Vol. 108, No. 6, pp. 1-25
7. The Role of the Teacher in the 21st Century	Journal of Educational Research	2010, Vol. 113, No. 7, pp. 1-30
8. The Impact of Teacher Education on Student Achievement	Journal of Educational Research	2015, Vol. 118, No. 8, pp. 1-35
9. The Role of the Teacher in the 21st Century	Journal of Educational Research	2020, Vol. 123, No. 9, pp. 1-40
10. The Impact of Teacher Education on Student Achievement	Journal of Educational Research	2025, Vol. 128, No. 10, pp. 1-45

RIKEN Mouse ESTs (Arakawa, T., et al. 2001)
Unpublished (2001)
On Nov 1, 1999 this sequence version replaced gi:6179369.
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
tel. 81-45-503-9222

FEATURES
source

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Location/Qualifiers
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/lab_host="DH10B"
/note="Site_1: Sali; Site_2: BamHI; cDNA library was
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Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in
RIKEN. Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA wa
primed with a primer [5',
GAGAGAGAGAGATCCCAAGAGCTCTTTTTTTTTTTTTTTTNN 3'], cDNA wa
prepared by using trehalose thermo-activated reverse
transcriptase and subsequently enriched for full-length
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to Rot = 10.0 and subtraction to Rot = 100.0. Second

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Ratio: 5.249
Percent Similarity: 98.558
Percent Identity: 98.558
Gaps: 0

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US-09-836-410A-1 x BE300741
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3 AAAATCTATAAGCATGCTGGAATATATAAAGAAGCTCAAGGTTGGATGGA 52
|||||
160 pGluAlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSerLysCysA 177
|||||
53 TGAGGCCACCGCCTTGACACACGACACAGATTTATCACTCCAATGTG 102
|||||
177 lalysTyrMetLeuLysAlaAsnLeuIleLysGluAlaGluMetCys 193
|||||
103 CAAATACATGCTTAAAGCCACCTGATTAAGAAGCTGAAGAATGTGC 152
|||||
194 SerLysPheThrArgGluGlyThrSerAlaValGluAsnLeuAsnGluMe 210
|||||
153 TCAAGTTTACAAGGGAAGGAACATCAGCGGTAGAGAATTTGAATGAAAT 202
|||||
210 tGlnCysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLysAlaMeta 227
|||||
203 GCACTGCATGTTGGTTCCTCAACAGAAATGTGCCAGCTTATAAAGCAATGA 252
|||||
227 snLysPheGlyGluAlaLeuLysLysCysHisGluIleGluArgHisPhe 243
|||||
253 ATAAATTTGGTGAAGCACTTAAAGAAATGTCAATGAGATTGAGAGACATTTT 302
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244 IleGluIleThrAspAspGlnPheAspPheHisThrTyrCysMetArgLy 260
|||||
303 ATAGAAATCACTGATGACCAAGTTTGACTTTCATCATCTACTGTATGAGGA 352
|||||
260 stleThrLeuArgSerTyrValAspLeuLeuLysLeuGluAspValLeuA 277
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353 GATTTACCTTAGATCATCATATGTGGACTTATTAACCTAGAAGATGTACTTC 402
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403 GACAGCATCCATTTTACTTCAAGGCGACGAAGAAATGCTATAGAGATCTAT 452
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294 LeuLysLeuHisAspAsnProLeuThrAspGluAsnLysGluHisGluAl 310
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453 TTGAAGCTTTCATGACCAACCCCTTACAGATGAGAATAAAGAACACGAAGC 502
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310 aAspThrAlaAsnMetSerAspLysGluLeuLysLysLeuArgAsnLysG 327
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503 TGATACAGCAACATGCTCTGACAAAGAGCTTAAGAAGCTAGTAAATAAAC 552
|||||
327 lnArgArgAlaGlnLysLysAlaGlnIleGluGluLysLysAla 343
|||||
553 AAAAGAAGAGCTCANAGAAAGAGCCCGAGATAGNAGAGAGAGAAAANAATGCA 602
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344 GluLysGluLysProGlnArgAsn 351
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602 CAAATCAAGGCAAGCAGAGAAAT 626

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seq name: qb_est1:AV227702

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LOCUS AV27702 988 bp mRNA linear EST 14-NOV-2001
DEFINITION AV27702 RIKEN full-length enriched, 14 days embryo liver Mus musculus cDNA sequence.
FEATURES
ORIGIN 1-988 ATCTGACGGTGTGTTTGATGAGGCGGTCG

ACCESSION	AV227702	GT:16385485
AV227702	2	

VERSION	KEYWORDS	EST.	hous.	MUS
SOURCE	ORGANISM			

Mus musculus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

Mon Jul 22 09:40:55 2002

us-09-836-410a-1.p2n.rst

Align seg 1/1 to: BJ057588 from: 1 to: 638

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173 nSerLysCysAlaLysTyrMetLeuLysAlaAsnLeuIleLysGluAlaG 190
|||||
65 TTCTAAATGTCAAAATATATGTTAAAGCAACCTGATTAAAGAGGCTG 114
|||||
190 lbcGluMetCysSerLysPheThrArgGluGlyThrSerAlaValGluAsn 206
|||||
115 AAGAAATGTCTCGAAATTTACAGGGAAGGACATCAGCAGTGGAAAT 164
|||||
207 LeuAsnGluMetGlnCysMetTrpPheGlnThrGluCysAlaGlnAlaTy 223
|||||
165 CTGAACGAGATGCGATGTCATGTGGTCCAGACAGAAATGTGCACAAGCTTA 214
|||||
223 rLysAlaMetAsnLysPheGlyGluAlaLeuLysCysHisGluIleG 240
|||||
215 CAAATCCATGAATAATATGCGCAGGCACTTAAATAATGCCATGAAATG 264
|||||
240 lueArgHisPheIleGluIleThrAspGlnPheAspPheHisThrTyr 256
|||||
265 AAGGCATTTGTAGAAATAACAGATGACCAGTTTGATTTCCACACTTAC 314
|||||
257 CysMetArgLysIleThrLeuArgSerTyrValAspLeuLysLeuG 273
|||||
315 TGTATGAGAAAAATTACACTCAGTCAATATGTGGACTTGTAAATTTAGA 364
|||||
273 uAspValLeuArgGlnHisProPheTyrPheLysAlaAlaArgIleAlaI 290
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365 GGATGTACTAAGCGCAGCATCCATTTTACTTCAAGGCTGCACGGATTGCAA 414
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323 uArgAsnLysGlnArgArgAlaGlnLysLysAlaGlnIleGluGluL 340
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615 GAAGATGATGATGAAGAGATTGGA 638
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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 19, 2002, 22:39:02 ; Search time 4131.3 Seconds
(without alignments)
17313.413 Million cell updates/sec

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Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1797656 seqs, 10463268293 residues

Total number of hits satisfying chosen parameters: 3595312

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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- 2: gb_htg.*
- 3: gb_in.*
- 4: gb_om.*
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- 23: em_pat.*
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- 25: em_pl.*
- 26: em_ro.*
- 27: em_sts.*
- 28: em_un.*
- 29: em_vl.*
- 30: em_htg_hum.*
- 31: em_htg_inv.*
- 32: em_htg_other.*
- 33: em_htgo_inv.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
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1.	3418	100.0	3418	6	AX285242	Sequence
2.	3418	100.0	3418	6	AX285294	Sequence
3	3418	100.0	3418	6	AX285296	Sequence
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7	1779	52.0	1779	6	AX285247	Sequence
c	1469.4	43.0	3324	5	AF247679	Xenopus l
8	1413	41.3	1413	6	AX285295	Sequence
10	1210.2	35.4	1802	9	AK023387	Sequence
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12.	1126	32.9	2859	9	AK001595	Rattus no
13	1079.6	31.6	1985	9	AK023402	Homo sapi
14	1038.4	30.4	145395	9	AC097376	Homo sapi
15	626.6	18.3	66729	2	AC102860	Mus muscu
c	617.8	18.1	710	6	AX210531	Sequence
17	448.2	13.1	843	9	BC017099	Homo sapi
18	439	12.8	2917	9	AB049844	Macaca fa
c	419	12.3	181573	2	AC020959	Mus muscu
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31	226	6.6	288	6	AX210600	Sequence
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ALIGNMENTS

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DEFINITION	AX285242	Sequence 1 from Patent WO0179506.				
ACCESSION	AX285242	Sequence 1 from Patent WO0179506.				
VERSION	AX285242.1	GI:17045930				
KEYWORDS	human.					
SOURCE	human.					
ORGANISM	Homo sapiens					
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.					
AUTHORS	1 (sites)					
TITLE	Gendron,R.L. and Paradis,H.					
JOURNAL	Treatment of Ocular neovascularization and related diseases					
FEATURES	Patent: WO 0179506-A 1 25-OCT-2001; Children's Hospital Research Foundation (US)					
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Query Match									
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Matches 3418; Conservative 0; Mismatches 0; Indels 0; Gaps 0;									
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ACCESSION AX285294
VERSION AX285294.1 GI:17045975
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SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (sites)
AUTHORS Gendron,R.L. and Paradis,H.
TITLE Inhibition of bone tumor formation using antisense cdna therapy
JOURNAL Patent: WO 0179505-A 2 25-OCT-2001;
CHILDREN'S HOSPITAL MEDICAL CENTER (US)
FEATURES Location/Qualifiers
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VERSION AX285296.1 GI:17045977
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (sites)
AUTHORS Gendron,R.L. and Paradis,H.
TITLE Inhibition of bone tumor formation using antisense cdna therapy
JOURNAL Patent: WO 0179505-A 4 25-OCT-2001;
CHILDREN'S HOSPITAL MEDICAL CENTER (US)
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 VERSION AF237622.1 GI:8164012
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 SOURCE Mus musculus
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 Gendron, R.L., Adams, L.C. and Paradis, H.
 Tubedown-1, A novel acetyltransferase associated with blood vessel
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 Dev. Dyn. 218 (2), 300-315 (2000)
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 Center, 3333 Burnet Avenue, Cincinnati, OH 45229-3039, USA
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RESULT 5
HSA314788
LOCUS
DEFINITION Homo sapiens mRNA for putative N-acetyltransferase.
ACCESSION AJ314788
VERSION AJ314788.1 GI:14589341
KEYWORDS N-acetyltransferase.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 5505)
AUTHORS Fluge,O., Bruland,O., Akslen,L.A., Varhaug,J.E. and Lillehaug,J.R.
TITLE Identification of NATH, a novel gene overexpressed in papillary
thyroid carcinomas
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 5505)
AUTHORS Fluge,O.
TITLE Direct Submission
JOURNAL Submitted (31-MAY-2001) Fluge O., Dept. of Molecular Biology,
University of Bergen, Thormohlens gt 55, N-5020 Bergen, NORWAY
COMMENT related entry AF327722.
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RESULT 7
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DEFINITION Sequence 6 from Patent WO0179506.
ACCESSION AX285247
VERSION AX285247.1 GI:17045931
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (sites)
AUTHORS Gendron, R.L. and Paradis, H.
TITLE Treatment of ocular neovascularization and related diseases
JOURNAL Patent: WO 0179506-A 6 25-OCT-2001;
Children's Hospital Research Foundation (US)
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AUTHORS		Expression of N-terminal acetyltransferase in Xenopus laevis		QY	565	tcttcaatcacttgagggtctttatatacagagataaagagaaggtggcaatcgtlagaagac	624
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REFERENCE		2 (bases 1 to 3324)		QY	625	tagtagtggttataaaactctctaaaagttgtcgctatttaaccccaatgatgatg	684
AUTHORS		Choi,S.-C., Kim,J. and Han,J.-K.		Db	1084	TAGTTGTTGGTTATGAACATCTCTGAAAGGCTGCCGTTTATTTAAACATGAATGATGATG	1143
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DEFINITION	Sequence 3 from Patent WO0179505.	1413 bp	DNA
ACCESSION	AX285295		
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SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
AUTHORS	1 (sites)		
TITLE	Gendron, R.L. and Paradis, H.		
JOURNAL	Inhibition of bone tumor formation using antisense cdna therapy		
FEATURES	Patent: WO 0179505-A 3 25-OCT-2001; CHILDREN'S HOSPITAL MEDICAL CENTER (US)		
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LOCUS			
DEFINITION	Homo sapiens cDNA FLJ13325 fis, clone OVARC1001762, weakly similar to N-TERMINAL ACETYLTRANSFERASE 1 (EC 2.3.1.88).		
ACCESSION	AK023387		
VERSION	AK023387.1		
KEYWORDS	oligo capping; fis (full insert sequence).		
SOURCE	Homo sapiens ovary, tumor tissue cDNA to mRNA, clone_lib:OVARC1		
ORGANISM	Homo sapiens		
REFERENCE	1 (sites)		
AUTHORS	Isogai,T., Ota,T., Hayashi,K., Sugiyama,T., Otsuki,T., Suzuki,Y., Nishikawa,T., Nagai,K., Sugano,S., Shiratori,A., Sudo,H., Wagatsuma,M., Hosoi,T., Kaku,Y., Kodaira,H., Kondo,H., Sugawara,M., Takahashi,M., Chiba,Y., Ishida,S., Murakawa,K., Ono,Y., Takiguchi,S., Watanabe,S., Kimura,K., Murakami,K., Ishii,S., Kawai,Y., Saito,K., Yamamoto,J., Wakamatsu,A., Nakamura,Y., Nagahara,K., Masuo,Y., Ninomiya,K. and Iwayanagi,T.		
TITLE	NEDO human cDNA sequencing project		
JOURNAL	Unpublished (2000)		
REFERENCE	2 (bases 1 to 1802)		
AUTHORS	Isogai,T. and Otsuki,T.		
TITLE	Direct Submission		
JOURNAL	Submitted (23-AUG-2000) to the DDBJ/EMBL/GenBank databases. Takao Isogai, Helix Research Institute, Genomics Laboratory; 1532-3 Yana, Kisarazu, Chiba 292-0812, Japan (E-mail:genomics@hri.co.jp, Tel:81-438-52-3951, Fax:81-438-52-3952)		
COMMENT	NEDO human cDNA sequencing project supported by Ministry of International Trade and Industry of Japan; cDNA full insert sequencing; Research Association for Biotechnology; cDNA library construction, 5'- & 3'-end one pass sequencing and clone selection; Helix Research Institute (supported by Japan Key Technology Center etc.) and Department of Virology, Institute of Medical Science, University of Tokyo.		
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McCarthy, M., McEwan, P., McKernan, K., McPheeters, R., Meldrim, J.,
Meneus, L., Mihova, T., Milenga, V., Murphy, T., Naylor, J., Nguyen, C.,
Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D.,
Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V.,
Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P.,
Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schuback, R.,
Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
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Topham, K., Travers, M., Travis, N., Trigliio, J., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.-J., Young, G.,
Zainoun, J., Zemzek, L., Zimmer, A. and Zody, M.
Direct Submission
Submitted (23-NOV-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: <http://www.seq.wi.mit.edu>
Contact: sequence-submissions@genome.wi.mit.edu
----- Project Information
Center project name: L19968
Center clone name: 511_H_12

* NOTE: This record contains 84 individual
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* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
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* 14183 14282: gap of 100 bp
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* 14283 14955: contig of 673 bp in length
* 14956 15055: gap of 100 bp
* 15056 15754: contig of 699 bp in length
* 15755 15854: gap of 100 bp
* 15855 16558: contig of 704 bp in length
* 16559 16658: gap of 100 bp
* 16659 17346: contig of 688 bp in length
* 17347 17446: gap of 100 bp
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* 18140 18239: gap of 100 bp
* 18240 18954: contig of 715 bp in length
* 18955 19054: gap of 100 bp
* 19055 19743: contig of 689 bp in length
* 19744 19843: gap of 100 bp
* 19844 20534: contig of 691 bp in length
* 20535 20634: gap of 100 bp
* 20635 21329: contig of 695 bp in length
* 21330 21439: gap of 100 bp
* 21439 22118: contig of 689 bp in length
* 22119 22218: gap of 100 bp
* 22219 22925: contig of 707 bp in length
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* 38114 38820: contig of 707 bp in length
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* 42019 42118: gap of 100 bp
* 42119 42838: contig of 720 bp in length
* 42839 42938: gap of 100 bp
* 42939 43639: contig of 701 bp in length

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OM nucleic - nucleic search, using sw model

Run on: July 20, 2002, 00:31:22 ; Search time 356.49 Seconds
(without alignments)
16461.650 Million cell updates/sec

Title: US-09-836-410A-2

Perfect score: 3418

Sequence: 1 caagtaacaccgcgaagtg.....atgcaataaaattgtttggg 3418

Scoring table:

IDENTITY_NUC

Gapop 10.0, Gapext 1.0

Searched: 1736436 seqs, 858457221 residues

Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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23: /SIDSI/gcgdata/hold-geneseq/geneseq-emb1/NA2001B.DAT:**
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	3418	100.0	3418	24	AAD22687
2	3418	100.0	3418	24	Human tubedown-1 c
3	3418	100.0	3418	24	Human tubedown-1 b
4	1502.2	43.9	2477	23	DNA encoding novel
5	1413	41.3	1413	24	Human tubedown-1 b
6	1210.2	35.4	1802	22	Human cDNA sequenc
7	1126	32.9	2859	22	Human cDNA sequenc
8	1079.6	31.6	1985	22	Human cDNA sequenc
9	617.8	18.1	710	22	Human differential

10	616.2	18.0	790	22	AAH06489	Human cDNA clone (
11	565	16.5	1225	20	AAH93092	Human cancer cell
12	555.6	16.3	781	20	AAZ15705	Human gene express
13	555.6	16.3	781	20	AAZ15705	Human validated ca
14	475.6	13.9	764	20	AAZ15983	Human gene express
15	475.6	13.9	764	20	AAZ15983	Human validated ca
16	398.4	11.7	774	20	AAH98889	Human validated ca
17	343.6	10.1	488	22	AAH12222	Human cDNA clone (
18	342.6	10.0	408	22	AAH12222	Novel human diagno
19	340	9.9	408	22	AAH12222	Novel human diagno
20	339.8	9.9	402	22	AAH12222	Novel human diagno
21	310.6	9.1	773	20	AAH93063	Human validated ca
22	281	8.2	404	22	AAH93063	Human validated ca
23	276.6	8.1	727	22	AAH93063	Human polynu
24	268	7.8	300	20	AAH93063	Human neuroblastom
25	258.4	7.6	300	20	AAH93063	Human cancer cell
26	258.4	7.6	300	20	AAH93063	Human gene express
27	242.6	7.1	399	20	AAH93063	Human cancer cell
28	239	7.0	255	22	AAH93063	EST clone CR392.
29	226	6.6	288	22	AAH93063	Rat differential t
30	215.4	6.3	255	22	AAH93063	Human differential t
31	214.4	6.3	300	20	AAH93063	Human cancer cell
32	212.8	6.2	300	20	AAH93063	Human cDNA clone (
33	188	5.5	653	22	AAH93063	Human gene express
34	163.8	4.8	802	21	AAH93063	Human colon cancer
35	149.4	4.4	297	21	AAH93063	Human secreted pro
36	73.4	2.1	2703	10	AAH93063	DNA encoding N-alp
37	73.4	2.1	2724	12	AAH93063	DNA encoding novel
38	57.2	1.7	654	23	AAH93063	DNA encoding novel
39	57.2	1.7	654	23	AAH93063	DNA encoding novel
40	57.2	1.7	654	23	AAH93063	DNA encoding novel
41	57	1.6	450	22	AAH93063	DNA encoding novel
42	56	1.6	887	22	AAH93063	Human reproductive
43	55.6	1.6	14006	24	AAH93063	Human neuroblastom
44	54.2	1.6	6171	24	AAH93063	Human immune syste
45	53.8	1.6	12007	24	AAH93063	Human immune syste

ALIGNMENTS

RESULT 1

AAD22687
ID AAD22687 standard; CDNA; 3418 BP.
XX
AC AAD22687;
XX
DT 26-FEB-2002 (first entry)
XX
DE Tubedown-1 (tbdn-1) protein encoding CDNA.
XX
KW Tubedown-1 protein; tbdn-1; ophthalmological; cytostatic; vulnerary;
KW cerebroprotective; angiogenesis inhibitor; ocular neovascularisation;
KW retinal disease; diabetic retinopathy; retinopathy of prematurity;
KW primary hyperplastic vitreous; macular degeneration; trauma; stroke;
KW haemorrhagic shock; arthritis; arteriosclerosis; delayed wound healing;
KW angiofibroma; granulation; nonunion fracture; retrolental fibroplasia;
KW solid tumour growth; chronic glaucoma; sickle cell retinopathy; cancer;
KW burn; scar; corneal neovascularisation; rubeosis iritis; uveitis;
KW gene therapy; ss.
XX
OS Unidentified.

Key	Location/Qualifiers
CDS	18..2189
FT	/*tag= a
FT	/product= "Tubedown-1 protein #2"
CDS	57..2189
FT	/*tag= b
FT	/product= "Tubedown-1 protein #3"
CDS	87..2189
FT	/*tag= c
FT	/product= "Tubedown-1 protein #4"

FT	CDS	408..2189	
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FT	FT	/product= "Tubedown-1 protein #1"	
FT	FT	408..2186	
FT	FT	misc_feature	
FT	FT	/note= "This region is specifically claimed as	
FT	FT	SEQ ID NO: 6 in claim 5 of the specification"	
XX	XX		
XX	XX		
XX	PN	WO200179506-A2.	
XX	PD		
XX	PD	25-OCT-2001.	
XX	XX		
XX	PF	17-APR-2001; 2001WO-US12548.	
XX	XX		
XX	PR	17-APR-2000; 2000US-197977P.	
XX	XX		
XX	PA	(CHIL-) CHILDRENS HOSPITAL RES FOUND.	
XX	XX		
XX	PI	Gendron RL, Paradis H;	
XX	XX		
XX	DR	WPI: 2002-026032/03.	
XX	DR	P-PSDB; AAEL13589, AAEL13590, AAEL13591, AAEL13592.	
PT	PT	Novel tubedown-1 protein comprising anti-angiogenic activity is useful	
PT	PT	for treating angiogenesis-associated disease related to ocular	
PT	PT	neovascularization, e.g., diabetic retinopathy, retinopathy of	
PT	PT	prematurity "	
XX	XX		
XX	PS	Claim 10; Page 56-58; 85pp; English.	
XX	XX	The present invention relates to tubedown-1 (tbdn-1) proteins and	
CC	CC	their corresponding cDNAs. Tbdn-1 proteins having anti-angiogenic	
CC	CC	activity are associated with acetyl transferase activity. They	
CC	CC	regulate endothelial differentiation through protein acetylation,	
CC	CC	DNA-binding or by interacting with and/or acetylating other protein	
CC	CC	targets important for endothelial differentiation. In normal adult	
CC	CC	eyes, tbdn-1 is expressed highly in the corneal endothelium proper	
CC	CC	and in the vascular endothelium of the limbus and retina. Tbdn-1	
CC	CC	proteins are useful for preventing, treating, inhibiting or delaying	
CC	CC	the onset of angiogenesis-associated disease related to ocular	
CC	CC	neovascularisation, e.g., a retinal disease, such as preferably	
CC	CC	diabetic retinopathy or retinopathy of prematurity, or primary	
CC	CC	hyperplastic vitreous, macular degeneration and any other conditions	
CC	CC	involving ocular neovascularisation. Tbdn-1 proteins are also useful	
CC	CC	for treating any pathological neovascularisation condition such as	
CC	CC	head trauma, spinal trauma, stroke, haemorrhagic shock, arthritis,	
CC	CC	arteriosclerosis, angiofibroma, delayed wound healing, granulations,	
CC	CC	cancer, burns, scars, nonunion fractures, retrolental fibroplasia,	
CC	CC	solid tumour growth. Proteins of the invention are also useful for	
CC	CC	treating ocular neovascularisation conditions such as chronic glaucoma	
CC	CC	sickle cell retinopathy, corneal neovascularisation, rubeosis iritis,	
CC	CC	uveitis, neovascularisation of the optic nerve. Sequences of the	
CC	CC	invention are also used in gene therapy. The present sequence is	
CC	CC	a cDNA encoding tubedown-1 proteins.	
XX	XX		
XX	SQ	Sequence 3418 BP: 1157 A; 604 C; 704 G; 953 T; 0 other;	
		Query Match 100.0%; Score 3418; DB 24; Length 3418;	
		Best Local Similarity 100.0%; Pred. No. 0;	
		Matches 3418; Conservative 0; Mismatches 0; Indels 0; Gaps	
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Qy	61	tttaccattttatagaagactatgaatggcgagcaaaaattttagaagaggtttaggaaaa 120	
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Qy	121	cacagcagacatctcctgataaagtggtattgaatagtgaactcctcttatcaga 180	
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Db 1361 cagcagaattgctcatgagctctattgagctctctgacacccctctgacagatgaga 1320
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ID	AAH77156 standard; cDNA; 3418 BP.
XX	AAH77156:
AC	21-JAN-2002 (first entry)
DT	Human tubedown-1 cDNA.
XX	Human; tubedown-1; tbdn-1; antisense; cytostatic; osteopathic;
KW	tubedown-1; osteosarcoma; Ewings sarcoma; metastasis; ss.
KW	Homo sapiens.
OS	Key Location/Qualifiers
FH	CDS 408..2189
FT	/tag= a
FT	/product= "tubedown-1"
XX	WO200179505-A2.
PX	25-OCT-2001.
PD	17-APR-2001; 2001WO-US12435.
PF	17-APR-2000; 2000US-197977P.
PR	17-APR-2001; 2001US-0836410.
XX	(CHIL-) CHILDRENS HOSPITAL RES FOUND.
PA	Gendron RL, Paradis H;
XX	WPI; 2002-017618/02.
PI	p-PSDB; AAG77907.
DR	Nucleic acid molecules antisense to the tubedown-1 gene prevent overexpression of tubedown-1 protein and are useful to treat osteosarcoma and Ewing's Sarcoma family of tumours - Claim 1; Page 36-38; 56pp; English.
XX	The sequence represents a new human gene, tubedown-1 (tbdn-1). The invention relates to a novel isolated nucleic acid of the tubedown-1 gene, and antisense nucleic acids to tbdn-1. The polynucleotides and protein of the invention have cytostatic and osteopathic activity. The polynucleotides of the invention may be used in antisense-therapy/gene therapy. They are useful in the treatment of bone tumours, especially osteosarcoma and Ewings sarcoma family of tumours. The compounds of the invention may also be useful for the prevention of metastases from these types of tumours, either alone or in combination with radiotherapy and/or chemotherapeutic agents.
CC	Sequence 3418 BP; 1157 A; 604 C; 704 G; 953 T; 0 other;
XX	Query Match 100.0%; Score 3418; DB 24; Length 3418;
SQ	Best Local Similarity 100.0%; Pred. No. 0;
	Matches 3418; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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DB	1 caagtaacaccgcgaagatgatagagatctgagagtcgagcatcgagcattgttgatgc 60
QY	61 ttaccattattagagactataaatggcagcaaaatttttagaagatttagaaaa 120
DB	61 ttaccattattagagactataaatggcagcaaaatttttagaagatttagaaaa 120
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QY	181 atcaagttcttcggaagcaggcttttatagagaagccctggaacacctttgtacctatg 240
DB	181 atcaagttcttcggaagcaggcttttatagagaagccctggaacacctttgtacctatg 240

QY 1321 acaagaacacgagcgtatcacagcaaaacatgtctgacaaagagctaaagaaactcgcta 1380
Db 1321 acaagaacacgagcgtatcacagcaaaacatgtctgacaaagagctaaagaaactcgcta 1380
QY 1381 ataaacaaagaagcgtcaaaaagaagccagatgtgaagaagagaaaaaaatgccgaaa 1440
Db 1381 ataaacaaagaagcgtcaaaaagaagccagatgtgaagaagagaaaaaaatgccgaaa 1440
QY 1441 aegaaagccgcgaacggaatccgaaaaaagaaaaagagatgatgacgaagaatacgag 1500
Db 1441 aegaaagccgcgaacggaatccgaaaaaagaaaaagagatgatgacgaagaatacgag 1500
QY 1501 gcccaaaaagaagcgtatccctgagaaaactgcccaggttgaaacctccattggaagaag 1560
Db 1501 gcccaaaaagaagcgtatccctgagaaaactgcccaggttgaaacctccattggaagaag 1560
QY 1561 ctattaaagtttttaacacacattgaagaacattggtgaagaacacagatagaactcatcttt 1620
Db 1561 ctattaaagtttttaacacacattgaagaacattggtgaagaacacagatagaactcatcttt 1620
QY 1621 ttgaccttgagatctacttttagaaaaaagtttctttttagtgcacaaatcagtaaaagc 1680
Db 1621 ttgaccttgagatctacttttagaaaaaagtttctttttagtgcacaaatcagtaaaagc 1680
QY 1681 gggcatttgcattgattctatgctatccctgcttcattgagtgcatgctgactctttc 1740
Db 1681 gggcatttgcattgattctatgctatccctgcttcattgagtgcatgctgactctttc 1740
QY 1741 attctgtgtgtaagtaagaagcttaccgaaacagtttagaacagattataaaacagaaa 1800
Db 1741 attctgtgtgtaagtaagaagcttaccgaaacagtttagaacagattataaaacagaaa 1800
QY 1801 tgaatgtcttttttgagcaacaaatccaaagaaattttaatgaaacctttctgaaagga 1860
Db 1801 tgaatgtcttttttgagcaacaaatccaaagaaattttaatgaaacctttctgaaagga 1860
QY 1861 attctgattcattgcacatagattatcagctgccaataatggtatattattagattctt 1920
Db 1861 attctgattcattgcacatagattatcagctgccaataatggtatattattagattctt 1920
QY 1921 ctatgcaaaaacgagcaaatagagctggcgacaacacttgatggatccctccacacagaa 1980
Db 1921 ctatgcaaaaacgagcaaatagagctggcgacaacacttgatggatccctccacacagaa 1980
QY 1981 accttcagacttgcattggaagtgttggagccttgtgtgagtgtagcctacgagactgta 2040
Db 1981 accttcagacttgcattggaagtgttggagccttgtgtgagtgtagcctacgagactgta 2040
QY 2041 aagaagctgcgcgaagcctcacagacaagtgttcataaagcttttcccttatgcttggctt 2100
Db 2041 aagaagctgcgcgaagcctcacagacaagtgttcataaagcttttcccttatgcttggctt 2100
QY 2101 tcatgctccttggatcacgaagatatagaagatcacagtgaacggagatagttctgcag 2160
Db 2101 tcatgctccttggatcacgaagatatagaagatcacagtgaacggagatagttctgcag 2160
QY 2161 aaacggaagaactggccaatgaaatctgaacatcattataaacaagcaaatggaatgacttt 2220
Db 2161 aaacggaagaactggccaatgaaatctgaacatcattataaacaagcaaatggaatgacttt 2220
QY 2221 ggaacatatctagtgataataattttgtcacgcacactgctgcattgctcttaactcac 2280
Db 2221 ggaacatatctagtgataataattttgtcacgcacactgctgcattgctcttaactcac 2280
QY 2281 agaattgagagagtaaatgttcttgccttcaaatagcttacctacgttttttacctgctgaa 2340
Db 2281 agaattgagagagtaaatgttcttgccttcaaatagcttacctacgttttttacctgctgaa 2340
QY 2341 aactatataaaatatctaacattacagatagattaggttcagtttcttaaaaaattaaa 2400
Db 2341 aactatataaaatatctaacattacagatagattaggttcagtttcttaaaaaattaaa 2400
QY 2401 gctgctaaaaattgagggttttaaaagaaaaaaaatccgtatctcttactcttccct 2460

Db 2401 gctgctaaaaattgagggttttaaaagaaaaaaaatccgtatccctatctccttccct 2460
QY 2461 tcccatgttttttaacttaatttataataaactctggaggtctataacagctaaactaagcagt 2520
Db 2461 tcccatgttttttaacttaatttataataaactctggaggtctataacagctaaactaagcagt 2520
QY 2521 gttgtggcagaataataacttttaatttcttctgtgagatgttctgctatatctcagacagca 2580
Db 2521 gttgtggcagaataataacttttaatttcttctgtgagatgttctgctatatctcagacagca 2580
QY 2581 taaaataaattgctgttttagcactgattctttcactgagcaacaagagttgttggggttt 2640
Db 2581 taaaataaattgctgttttagcactgattctttcactgagcaacaagagttgttggggttt 2640
QY 2641 tagcatctgcctgattctgttacgggtgttggtgattgaccataggaagtatgcaatgcta 2700
Db 2641 tagcatctgcctgattctgttacgggtgttggtgattgaccataggaagtatgcaatgcta 2700
QY 2701 atcactgtgtacagagccgtctacaaacatgcttgacgtgttagagactgggacacata 2760
Db 2701 atcactgtgtacagagccgtctacaaacatgcttgacgtgttagagactgggacacata 2760
QY 2761 gctacaaagcggatttaagtgaacctagaaaggtgttcaagtaacgtgtgtgtgtttccaaa 2820
Db 2761 gctacaaagcggatttaagtgaacctagaaaggtgttcaagtaacgtgtgtgtgtttccaaa 2820
QY 2821 attcactgtacatgatcagtttgggtgttctgttacacagtttttaaccggaagacacag 2880
Db 2821 attcactgtacatgatcagtttgggtgttctgttacacagtttttaaccggaagacacag 2880
QY 2881 ttggaacaattctcaatttaactaaacttgaaacttaaaataaactgcaaaactttat 2940
Db 2881 ttggaacaattctcaatttaactaaacttgaaacttaaaataaactgcaaaactttat 2940
QY 2941 cattgttttggccaaaactgtttaaactgttaactgcaagaacaaactcactgtgatgtg 3000
Db 2941 cattgttttggccaaaactgtttaaactgttaactgcaagaacaaactcactgtgatgtg 3000
QY 3001 caccacataattatgcaagcatgaattttcacctgagagtgaagaaagaaactctacc 3060
Db 3001 caccacataattatgcaagcatgaattttcacctgagagtgaagaaagaaactctacc 3060
QY 3061 atggcttgaagtacagagcagaactcctgactacacattctctatgactgatgaagagact 3120
Db 3061 atggcttgaagtacagagcagaactcctgactacacattctctatgactgatgaagagact 3120
QY 3121 aatatctaaaacccctcagcagcctgttcccgatatacgagaaaaagtgctgcagttttaga 3180
Db 3121 aatatctaaaacccctcagcagcctgttcccgatatacgagaaaaagtgctgcagttttaga 3180
QY 3181 taaccttgggaacttttccacagtgctacaggtttgttaaacttgaagcccttcttct 3240
Db 3181 taaccttgggaacttttccacagtgctacaggtttgttaaacttgaagcccttcttct 3240
QY 3241 aagaataataattctcgcctcagttgtttcaggaagccagacatttctgtaatttttaag 3300
Db 3241 aagaataataattctcgcctcagttgtttcaggaagccagacatttctgtaatttttaag 3300
QY 3301 gcccaagatttttttcaataacagacagcagcttttttccctgcagtttcaaaatgtaat 3360
Db 3301 gcccaagatttttttcaataacagacagcagcttttttccctgcagtttcaaaatgtaat 3360
QY 3361 ttctttttttttttttgttgcataaaggtacacaaatgcaaatgcaaatgttttggg 3418
Db 3361 ttctttttttttttttgttgcataaaggtacacaaatgcaaatgcaaatgttttggg 3418

RESULT 3
AAH77158/C
ID AAH77158 standard; cDNA; 3418 BP.
XX
AC AAH77158;

XX	DT	21-JAN-2002	(first entry)	
XX	DE	Human tubedown-1	base pairs 3418-1 antisense cDNA.	
XX	DE	Human: tubedown-1; tbdn-1; antisense; cytostatic; osteopathic; bone tumour; osteosarcoma; Ewings sarcoma; metastasis; ss.		
XX	KW	Homo sapiens.		
OS	XX	WO2001179505-A2.		
PN	XX	25-OCT-2001.		
PD	XX	17-APR-2001; 2001WO-US12435.		
PF	XX	17-APR-2000; 2000US-197977P.		
XX	PR	17-APR-2001; 2001US-0836410.		
XX	XX	(CHIL-) CHILDRENS HOSPITAL RES FOUND.		
XX	PA	Gendron RL, Paradis H;		
XX	PI	WPI; 2002-017618/02.		
XX	DR	Nucleic acid molecules antisense to the tubedown-1 gene prevent overexpression of tubedown-1 protein and are useful to treat osteosarcoma and Ewing's Sarcoma family of tumours -		
XX	PT	Claim 7; Page 39-41; 56pp; English.		
XX	PS	The sequence represents tubedown-1 (tbdn-1) bases 3418-1 antisense cDNA. The invention relates to a novel isolated nucleic acid of the tubedown-1 gene, and antisense nucleic acids to tbdn-1. The polynucleotides and protein of the invention have cytostatic and osteopathic activity. The polynucleotides of the invention may be used in antisense-therapy/gene therapy. They are useful in the treatment of bone tumours, especially osteosarcoma and Ewings sarcoma family of tumours. The compounds of the invention may also be useful for the prevention of metastases from these types of tumours, either alone or in combination with radiotherapy and/or chemotherapeutic agents.		
XX	CC	Sequence 3418 BP; 953 A; 704 C; 604 G; 1157 T; 0 other;		
XX	SQ	Query Match 100.0%; Score 3418; DB 24; Length 3418; Best Local Similarity 100.0%; Pred. No. 0; Matches 3418; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		
QY	1	caagtaacacccccaagatgataagagatctgcagagtgagcatcatcgattggtatgc 60		
DB	3418	CAAGTACACCCCAAGATGATAGAGATCTGCAGATGAGCATCATGATTGTTATGC 3359		
QY	61	ttaccattattagaagactgaaatggcagcaaaaattttagaagaagtttaggaaaa 120		
DB	3358	TTTACCATTATTAGAGACATATGAATGGCAGCAAAATTTTAGAAGAGTTTAGGAAA 3299		
QY	121	cacagcagacatctcttgataaagtggattgaatatagtgaaactcctcttatcaga 180		
DB	3298	CACAGCAGACATCTCTGTATAAAGTGGATTATCAATATAGTGAACCTCTTATATCAGA 3239		
QY	181	atcaagttcttcgggaagcagggtctttattagagaagccctggaaacattcttgacctatg 240		
DB	3238	ATCAAGTCTTCTCGGGAAGCAGGCTCTTATAGAGAAGCCCTGGAAACATCTTTGTACCTATG 3179		
QY	241	aaaagcagatttgtataaactgctgtgtgaagaacccaagggaacctctctgttcagtt 300		
DB	3178	AAAGCAGATTGTGTAATACTTGTCTGCTCAAGAAACCAAGGGGAACCTTCTGTTCAGT 3119		
QY	301	tgtgtcgtttggaagatgctgctgaagctttatagagattacaaagagaggaatcctgaaa 360		
DB	3118	TCGTGCGTTTCGAAAGATGCTGCTGAGCTTTATAGAGGATTACAAGAGAGGAATCTGAAA 3059		

XX WO200175067-A2.
 XX 11-OCT-2001.
 XX 30-MAR-2001; 2001WO-US08631.
 XX 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX (HYSE-) HYSEQ INC.
 PA Drmanac RT, Liu C, Tang YT;
 PI WPI: 2001-639362/73.
 DR P-PSDB; ABG07738.
 XX
 XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.
 XX
 XX Claim 1; SEQ ID NO 7729; 103pp; English.
 XX The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 XX Sequence 2477 BP; 859 A; 437 C; 522 G; 659 T; 0 other;

Query Match 43.9%; Score 1502.2; DB 23; Length 2477;
 Best Local Similarity 80.7%; Pred. No. 0;
 Matches 2082; Conservative 0; Mismatches 218; Indels 279; Gaps 17;

Qy 322 ctgacgtttatagaggattacaagagggaatcctgaaaattgggacctattacaaggct 381
 Db |||||
 Qy 449 -cagatgtttatagaggattgcaagagagaatcctgaaaactggcctattacaaggct 508
 Db |||||
 Qy 382 tagaaaaagcactgaagccagcctaattgttagaacggctaaaaatatagagaagcct 441
 Db |||||
 Qy 509 tggaaaaagcactcaagccagcctaattgttagaacggctaaaaatatagagaagcct 568
 Db |||||
 Qy 442 ggactaaataccaggggactcgtccaagaagctgccttaaaactttttatctggag 501
 Db |||||
 Qy 569 ggactaaataccaggggactcgtccaagaagctgccttaaaactttttatctggag 628
 Db |||||
 Qy 502 agaaqtttaaggagtggttgatagggttcctaagatgaattcagcaagggtgtccac 561
 Db |||||
 Qy 629 agaagtttaagaatgtttgataagttcctaagatgaattcagcaagggtgtccac 688
 Db |||||
 Qy 562 ctgttccaatactctgaggtcttta-tacagagataagagaag-gtggcaactgtaga 619
 Db |||||
 Qy 689 cagttccaatactttaagatcattactaccaagacaagaaggtgtggcaactataga 748
 Db |||||
 Qy 620 agaactagtagtggttatgaaactctctaaaaagttgtgcctatttaaccccaatga 679
 Db |||||
 Qy 749 aaagttagtagtggttatgaaactctctaaaaagctgcggctatttaaccccaatga 808
 Db |||||
 Qy 680 tgatgaaaggaggaaacctccaaccacattactttgggtccagtactatttggcacagca 739
 Db |||||
 Qy 809 tgatgaaaggaggaaacctccaaccacattactttgggtccagtactatttggcacagca 868
 Db |||||
 Qy 740 ttatgataaaattgtcagccatccattctgtgaaatacataataacttgcgaattgaag 799
 Db |||||
 Qy 869 ttatgataaaattgtcagccatccattctgtgaaatacataataacttgcgaattgaag 928
 Db |||||
 Qy 800 tacaccaacttgatagaactctttgtgaaagctaaaaactataaactgcgtgggaa 859
 Db |||||
 Qy 929 tacacctataatagaactctttctgtgaaagctaaaaactataaactgcgtgggaa 988
 Db |||||
 Qy 860 tattaagaagctgcagtgatgaagccagccctggacacagcagacagatt 919
 Db |||||
 Qy 989 tattaagaagctgcagtgatgaagccagccctggacacagcagacagatt 1048
 Db |||||
 Qy 920 tattaattccaagtgcaaaatacatgtttaaagccaacctgatttaaagagggtgaaga 979
 Db |||||
 Qy 1049 tatcaactccaatgfcgaaaatacatgtctaaagccaacctgatttaaagagggtgaaga 1108
 Db |||||
 Qy 980 aatgtgtccaagttcacgagggaagaaacttcagcgttagagaacctgaatgaatgca 1039
 Db |||||
 Qy 1109 catgagctcaagtttacaagggggacatcagcggtagagaatttgaatgaattca 1168
 Db |||||
 Qy 1040 gtatgtgtgttcagacagagtgctcagcgatatacaaaagcaatgaacaaatttggta 1099
 Db |||||
 Qy 1169 gtcatgtgtgttcacatacagaagtgtcccagcttataaagcaatgaataatttggta 1228
 Db |||||
 Qy 1100 agcacttaagaatgtcatgaattgagagacattttatagaataaccagatcacagctt 1159
 Db |||||
 Qy 1229 agcacttcagaatgtcatgagattgagagacattttatagaataaccagatcacagctt 1288
 Db |||||
 Qy 1160 tgactttcatatactgtatgaggaagatcaccttagatcatatgtggaactattaaa 1219
 Db |||||
 Qy 1289 tgactttcatatactgtatgaggaagatttaccttagatcatatgtggaactattaaa 1348
 Db |||||
 Qy 1220 actagaagatgtacttcgacagcaaccttttacttctcaag--cagcgagaattgtctatt 1277
 Db |||||
 Qy 1349 actagaagatgtacttcgacagcaaccttttacttcaagggcgaggaagaattgtctata 1408
 Db |||||
 Qy 1278 gagatctattttaa-gtttcatacaacctctgcagagatgagacaagaacacagggc 1336
 Db |||||
 Qy 1409 gagatctatttgaaggcttcatagacaacctcttcacagatgaggaattaaaggaaacgg 1468
 Db |||||
 Qy 1337 tg-----atacagcaaacatgtctgacaagagctcaaaagactgcgtataaacaagaag 1390
 Db |||||
 Qy 1469 aggtgtgatacagccaaacatgtctgacaagagctcaaaagagctacgtataaacaagaag 1528
 Db |||||
 Qy 1391 aagagctcaaaagaaagcccgatttgaagaagagaaaaaaatccgaaaaaagaagcc 1450
 Db |||||

Db	153	CAGCGAATTTGTTATTTGAGATCTATTTGAAGCTTCATGACAACTCTGACAGATGAGA	94
Qy	1321	acaaagacacgagctgtatcacgcaaacatgtctgacaaagagctaaagaacactgcgta	1380
Db	93	ACAAAGAACACGAGGCTGTATACAGCAAAACATGTCTGACAAAGAGCTTAAAGAAACTGCCTA	34
Qy	1381	ataacaaagaagagctcaaaagaagccaga	1413
Db	33	ATAAACAAAGAAGAGCTCAAAAGAAAGCCCCAGA	1

AAH16408
ID AAH16408 standard: cDNA: 1802 BP.

ID AHH10400 Standard, SEMI, 11-1
 XX
 AHH10400

AC AAH164087

XX
DT 26-JUN-2001 (first entry)

XX
DE Human cDNA sequence SEQ ID NO:15380

Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.

[illegible]

XX
Homo sapiens

PN EP1074617-A2
XX

PD 07-FEB-2001.
VY

28-JUL-2000;

XX
PR 29-JUL-1999;

PR 27-AUG-1999;
PR 11-JAN-2000;

PR 02-MAY-2000;
PP 09-JUN-2000;

XX
05 DEC 2007

XX
PA (HELI-) HELI

PI Ota T, Isog
PI Ishii S, Su

XX WPI: 2001-31
DRXX
Primer sets

PI	PImer	DCB	full-length
PT			

PT and/or diagnosis
PT full-length

XX
PS Claim 8; SEQ

XX The present

CC full-length

CC comprises: (1)
CC to the compl

CC the 5602 nucleotides

cc of an oligonucleotide complementary to the 3' end of the mRNA.

CC complement sequence and
CC sequence and
CC sequence and

CC polynucleotides

CC oligonucleotides

CC the 5'-end
CC the specific

CC in gene the
CC particularly

CC detection a
CC the full-le

CC The full re
CC cDNAs easil

CC	AAH13633	to
CC	AAB95893	re

cc represent o

CC of the present invention.

XX Sequence 1802 BP; 644 A; 298 C; 395 G; 465 T; 0 other;
SQ

Query Match 35.4%; Score 1210.2; DB 22; Length 1802;
Best Local Similarity 92.3%; Pred. No. 1.2e-271;
Matches 1287; Conservative 0; Mismatches 103; Indels 5; Gaps 1;

```

QY 27 gatctcagagtgagcatcatcgattggttatgct-----ttaccattttattagaagct 81
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Db 408 gaactgcgagagagcatcatcgattggttatgctattgcttaccattttattagaagct 467
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QY 82 atgaatggcagcaaaatttttagaagagtttagaagaaacacagcagacatctctctgata 141
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 468 atgaatggcagcaaaagatttttagaagaaatttagaagaaacacagacatcccttgaca 527
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QY 142 aagtgattatgaatatagtaactctcttatatcagaatcaagttcttcgggaagcag 201
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Db 528 aagtgattatgaatatagtaactctcttatatcagaatcaagttcttcgggaagcag 587
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QY 202 gtctttatagaagcccttggaacatctttgtacatgaaagcagatttgtgataaac 261
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 588 gtctctatagaagccttgggaacatctttgtacatgaaagcagatttgtgataaac 647
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 262 ttgctgttgaagaacacaaaggggaactctctgttcagttgtgtctgttgggaagctg 321
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Db 648 ttgctgtgaagaacacaaaggggaactctctgttcagttgtgtctgttgggaagctg 707
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 322 ctgagctttatagaggttacagagaggaatcctgaaattgggcctattacaaagct 381
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 708 cagatgtttatagaggttgcaagagagaaatcctgaaactgggcctattacaaagct 767
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 382 tagaanaagcactgaagcagcctaataatgttagaagcgttaaaaataataggaagcct 441
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
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QY 442 ggaactaaatccccggggaactctgccaagaagctcccttaaacctttttatctcgag 501
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 828 agactaaataccccggggaactctgccaagaagctcccttaaacctttttatctcgag 587
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 502 agaaagtttaagagtggtttgtagaggttccctaaagatgaatttcagcaagggctgtccac 561
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Db 898 agaaagtttaagagtggtttgtagaggttccctaaagatgaatttcagcaagggctgtccac 947
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
QY 562 ctgtcttcaatcccttgaggtcttttacagacagataaagagaagtggaatctgtagaag 621
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 948 cagcttcaatcccttgaggtcttttacagacagataaagagaagtggaatctgtagaag 1007
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QY 622 aactagatgttggttatgaaactctcttaaaaagttgtcgccctatttaacccccaatgatg 681
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Db 1008 agttagtagtaggttatgaaacctctcttaaaaagctgccggttatttaacccccaatgatg 1067
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QY 682 atggaagaggggaacccctcccaaccacattactttgggtccagactattttggcagacatt 741
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RESULT 7

AAH14477

ID AAH14477 standard; cDNA; 2859 BP.

XX AC AAH14477;

XX DT 26-JUN-2001 (first entry)

XX DE Human cDNA sequence SEQ ID NO:11977.

XX KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.

XX OS Homo sapiens.

XX PN EP1074617-A2.

XX PD 07-FEB-2001.

XX PF 28-JUL-2000; 2000EP-0116126.

XX PR 29-JUL-1999; 99JP-0248036.

XX PR 27-AUG-1999; 99JP-0300253.

XX PR 11-JAN-2000; 2000JP-0118776.

XX PR 02-MAY-2000; 2000JP-0183767.

XX PR 09-JUN-2000; 2000JP-0241899.

XX PA (HELI-) HELIX RES INST.

XX PI Ota T, Isogai T, Nishikawa T, Hayashi K, Salto K, Yamamoto J;

XX PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

XX WPI; 2001-318749/34.

XX PT Primer sets for synthesizing polynucleotides, particularly the 5602

XX PT full-length cDNAs defined in the specification, and for the detection

XX PT and/or diagnosis of the abnormality of the proteins encoded by the

XX PT full-length cDNAs -

XX PS Claim 8; SEQ ID 11977; 2537pp + CD ROW; English.

XX

The present invention describes primer sets for synthesising 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesising polynucleotides particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAH03166 to AAH13628 and AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632 represent oligonucleotides, all of which are used in the exemplification of the present invention.

Sequence 2859 BP: 921 A: 476 C: 547 G: 915 T: 0 other;
XX
XX

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Best Local Similarity	90.2%	Pred. No. 5.6e-252;		
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Qy	1942	agctggcgacaaacttgatggtatccctcaccacgaagaaccttcagactgcgatggaag	2001
Db	61	agttggcaacacacttgatgaa tctctcatcacagaaacctccagacactgatatggaag	120
Qy	2002	tgttgggaagccttggatggtgagctcacgagactgtaaagaagctgccagaagcctaca	2061
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Qy	2182	aaactcgaacatcattaaacgaacaaatggatgactttggaccatactagtgtataat	2241
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Qy	2242	attttgtacgcacactgcgtgcatgctcttacttacacagaatcagagaggtaaatgtt	2301
Db	361	attttgtacgcacactgcgtgcatgctcttacttacacagaatcagagaggtaaatgtt	420
Qy	2302	cttgctctcaaat--gtcttcagcttttttattcctgctg--aaactatataaaaaatc	2358
Db	421	cttgctctcaaatggttttcagcttttttattcctgctgaaaaagtatataaaaaatc	480
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Db	481	taacattacagatataggtttcagttctttaaaaaatt--aaagctgctcaaaattgagtg	539
Qy	2419	tttaaaagaaaaaaatccgtatacttacttcacttcctccacttcctccactgtttttaactaa	2478
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Qy	2479	tttatataaaactcgagcgtataacagcttaacagcttaacatgacaggtgtgtggcagaataattac	2538
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DEPT. T 8

RESULTS	6
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ID	AAH16424 standard; cDNA; 1985 BP.
XX	
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AC	AAH16424;
XX	
DT	26-JUN-2001 (first entry)
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DE	Human cDNA sequence SEQ ID NO:15407.
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XX	Humao. primer. detection; diagnosis; antisense therapy; gene therapy; ss

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2724	Y	caacacatgcttgacgtttgtagagactgggacacatagctaccacagc-ggatttaagtaa	2782
120	b	caacacatgcttgatgtgttagaaactggacatatagataccaagcaaaattataagaa	179
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2843	Y	gggtgttctgtacacagtttttaaacggaagcaactgttggaacaaactcaa-tttaac	2901
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297	b	taaaactggaagaactaaaaacaaatgcacaaacctttcagcattgttttgccaaaacttgt	356
2962	Y	taaaactgtaatgcagaaccaaatagcactgtgagtgtggcaccacactaaattatgcaagca	3021
357	b	taaaactgtaatgcagaaccaaatagcactgtgagtgtggcaccacactaaata-gcaagca	415
3022	Y	tgaatttttccacctgaagtgaaaaaagaaaactctaccatggcttgaagttaacagagc	3081
416	b	tgaatttttccccaagagtgtaaaaaagaaaaactaccatgtgcttgaagtta-aagagc	474
3082	Y	agaacctctgactaccattctactgactgatgaagagactaata-tctaaaaacctcagcag	3140
475	b	agaacctctgactaccattctactgactgatcaaaagactaatagttaaaaaacctcagcag	534
3141	Y	gccttgttccagatgcag--aaaaagtctgcagtttagataacctctgggaacttt	3197
535	b	gccttgttccagatgcagaaaaaagtgctgcagtttagataacctct-aggaaattttt	593

can be used

The present invention describes a library of human polynucleotides comprising the sequences given in AAZ12532 to AAZ17779. Also described is a method of detecting differentially expressed genes correlated with the cancerous state of a mammalian cell, comprising detecting at least one differentially expressed gene product in a test sample from a cell suspected of being cancerous, where the gene product is encoded by one of the 5248 polynucleotide sequences given in AAZ12532 to AAZ17779. The polynucleotides can be used as a source of primers and probes, which can be used for a variety of purpose, e.g. detection of expression levels, mapping, tissue typing or profiling, forensics, genetic analysis and detection of polymorphisms. Polypeptides encoded by the polynucleotides can be used for raising antibodies for experimental, diagnostic and

therapeutic purposes. The polynucleotides may also be used to construct arrays for diagnostics (which may be used to determine function of an encoded protein); and to detect differences in expression levels between two cells (e.g. to identify abnormal or diseased tissue in a human, to identify a genetic predisposition or susceptibility to a disease such as cancer). The polynucleotides of the invention are especially used in the diagnosis, prognosis and management of colorectal cancer, breast cancer, and lung cancer. The polynucleotides can also be used to screen for peptide analogues and antagonists.

Sequence 781 BP; 263 A; 140 C; 156 G; 205 T; 17 other:

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597	QY	aaagagaagtgycaatcgtagaagaactagtagtctggttatgaacctctctaaaaagt	656
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657	QY	tgctgcctatttaaccccaatgatgtgaaagaggaacctccaacacattactttgg	716
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777	QY	ttacataaatactgcaattggaagtacacacacattgtatagaactcttcttctgtataaagct	836
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RESULT 13
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AAX99053;
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DT 24-SEP-1999 (first entry)

XX DE XX KW KW KW KW KW KW KW KW OS XX PN XX PD XX PF XX PR PR PR PR PA PA XX PI PI PI PI XX DR XX PT XX PS XX CC

Human validated cancer cell derived cDNA #375.

Cancer; human; colon; breast; lung; transmembrane receptor; ATPase; integral membrane protein; aspartyl protease; GATA family; wnt family; transcription factor; G-protein alpha subunit; protein phosphatase; cholesterol binding protein; diacylglycerol binding protein; trypsin; protein kinase; tyrosine phosphatase; developmental signalling protein; WW/rsp5/WWP domain; therapy; forensic; genetic mapping; diagnostic; detection; treatment; cervical; melanoma; colorectal adenocarcinoma; Wilms' tumour; retinoblastoma; sarcoma; myosarcoma; lung carcinoma; leukemia; lymphoma; dysplasia; hyperplasia; endometrium; adrenal; prostate; ss.

Homo sapiens.

WO9933982-A2.

08-JUL-1999.

22-DEC-1998; 98WO-US27610.

21-DEC-1998; 98US-0217471.

23-DEC-1997; 97US-0068755.
03-APR-1998; 98US-0080664.

21-OCT-1998; 98US-0105234.
27-OCT-1998; 98US-0105877

(CHTP \ CHTRON CORP.

(HYSE-) HYSEQ INC.

Crkvenjakov R, Dickson M, Drmanac R, Drmanac S;

Jones LW, Kassam A, Kennedy GC, Kita D, Labat I;

Stache-Crain B, Sudduth-Klinger J, Williams LT;

WPI; 1999-430243/36.

New isolated human polynucleotides

Claim 1; Page 564; 591pp; English.

This invention describes novel isolated human polynucleotides obtained by screening for differential expression in colon cancer, breast cancer and lung cancer cell lines. The polynucleotides of the invention are represented in AX98275-X9918 and encode polypeptides of protein families selected from 4 transmembrane segments integral membrane proteins, 7 transmembrane receptors, ATPases associated with various cellular activities (AAA), eukaryotic aspartyl proteases, GATA family of transcription factors, G-protein alpha subunit, phospholipases or diacylglycerol binding proteins, protein kinase, protein phosphatase 2C, protein tyrosine phosphatase, trypsin, wnt family of developmental signalling proteins and WW/rsp5/WWP domain containing proteins. The encoded polypeptides also have a functional domain selected from Ank repeat, basic region plus leucine zipper transcription factors, bromodomain, EF-hand, SH3 domain, WD domain/G-beta repeats, zinc finger (C2H2 type), zinc finger (CCHC class), and zinc-binding metalloprotease domain. The polynucleotides encode polypeptides with similarity to known protein families and are predicted to have similar properties. The novel polynucleotides can be used to develop products for use as therapeutic agents and in forensics, genetic analysis, mapping and diagnostic applications. In particular, the product can be used for the detection and management of cancers. They can be used for treating e.g. cervical cancers, melanomas, colorectal adenocarcinomas, Wilms' tumour, sarcomas, retinoblastoma, myosarcomas, lung carcinomas, leukemias, such as chronic myelogenous leukemia, promyelocytic leukemia, monocytic leukemia, and myeloid leukemia, and lymphomas such as histiocytic lymphoma, anhydric hereditary ectodermal dysplasia, congenital alveolar dysplasia, epithelial dysplasia of the cervix, fibrous dysplasia of bone, and mammary dysplasia, hyperplasias, e.g. endometrial, adrenal, breast, prostate or thyroid hyperplasias or pseudoepitheliomatous hyperplasia of the skin.

[illegible]

Mon Jul 22 09:40:57 2002

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QY 1129 gacattttatagaaatcacccgatga-ccagtttgactttcatatactactgtatga-ggaa 1186
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QY 1187 gatcaccccttag 1198
Db 665 nattaaccccttag 676
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Search completed: July 20, 2002, 02:41:40
Job time: 7818 sec

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gb_pat:AX137735	+	37.00	AX137735	7.2e+04	36	AX137735	Sequence 8 from Pate
gb_pat:AX207592	+	37.00	75.85	7.2e+04	36	AX207592	Sequence 1 from Pate
gb_pat:AR004318	+	37.00	75.61	7.5e+04	37	AR004318	Sequence 1 from Pate

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KEYWORDS			
SOURCE	synthetic construct.		
ORGANISM	synthetic construct		
	artificial sequence.		
REFERENCE	1 (bases 1 to 49).		
AUTHORS	Hogrefe,H.H.; Cline,J.M., Hansen,C.J. and Borns,M.C.		
TITLE			

JOURNAL Patent: WO 0109347-A 34 08-FEB-2001;

STRATAGENE (US)

[illegible]

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  VERSION AX135674.1 GI:14271944
  KEYWORDS :
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  ORGANISM artificial sequence.
  REFERENCE 1 (bases 1 to 49)
  AUTHORS Sorge,J.A.
  TITLE Methods for detection of a target nucleic acid sequence
  JOURNAL Patent: WO 0132922-A 15 10-MAY-2001;
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ACCESSION AR126194		
VERSION AR126194.1 GI:14112787		
KEYWORDS	Unknown.	
SOURCE	Unknown.	
ORGANISM	Unclassified.	
REFERENCE 1 (bases 1 to 45)		
AUTHORS Ruoslahti, E. and Pasqualini, R.		
TITLE NGR receptor and methods of identifying tumor homing molecules that		
JOURNAL home to angiogenic vasculature using same		
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AUTHORS Silke, N., Lerch, K. and Muheim, A.		
TITLE Cloning, expression and production of tasty peptides		
JOURNAL Patent: EP 0832972-A 11 01-APR-1998;		
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  ACCESSION AX221610
  VERSION AX221610.1 GI:15549334
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    artificial sequence.
  REFERENCE
    1 (bases 1 to 48)
  AUTHORS
    Blatt, L., McSwiggen, J. and Chowrira, B. M.
  TITLE
    Method and reagent for the modulation and diagnosis of cd20 and
    nogo gene expression
  JOURNAL
    Patent: WO 0159103-A 7052 16-AUG-2001;
    RIBOZYME PHARMACEUTICALS, INC. (US); Blatt, Lawrence (US);
    McSwiggen, James (US); Chowrira, Bharat M. (US)
FEATURES
  source
    Location/Qualifiers
      1..48
        /organism="synthetic construct"
        /db_xref="taxon:32630"
        /note="Nucleic Acid"
BASE COUNT
ORIGIN

alignment_scores:
  Quality: 40.50      Length: 19
  Ratio: 2.893       Gaps: 1
Percent Similarity: 73.684 Percent Identity: 47.368

alignment_block:
  US-09-836-410A-1 x AX221610 ..
  Align seg 1/1 to: AX221610 from: 1 to: 48

381 ProLeuGluAlaIleLysPheLeuThrProLeuLysAsnLeuVally 397
4 CCCTTGAGGAA.....ACTCCCTCAAGGACATCGTCCG 38
397 sAsnLys 399
39 GGATAAA 45
seq_name: gb_pr:HUMRPY59

seq_documentation_block:
  LOCUS HUMRPY59 48 bp mRNA linear PRI 27-SEP-2001
  DEFINITION Homo sapiens mRNA for ribosomal protein L24, partial cds.
  ACCESSION D28400
```

```
VERSION D28400.1 GI:461272
KEYWORDS
SOURCE
  Homo sapiens lymphoma cell_line:U937 cDNA to mRNA,
  clone_lib:U937/pkai clone:HP00302.
ORGANISM
  Homo sapiens
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
  1 (bases 1 to 48)
  AUTHORS
    Kato, S., Sekine, S., Oh, S. W., Kim, N. S., Umezawa, Y., Abe, N.,
    Yokoyama-Kobayashi, M. and Aoki, T.
  TITLE
    Construction of a human full-length cDNA bank
  JOURNAL
    Gens. 150 (2), 243-250 (1994)
  MEDLINE
    95121910
REFERENCE
  2 (bases 1 to 48)
  AUTHORS
    Kato, S.
  TITLE
    Direct Submission
  JOURNAL
    Submitted (03-FEB-1994) Seishi Kato, Research Institute of National
    Rehabilitation Center for the Disabled, Department of
    Rehabilitation Engineering; 4-1 Namiki, Tokorozawa, Saitama
    359-8555, Japan (E-mail:seishi@rehab.go.jp,
    Tel:81-42-995-3100(ex.2568), Fax:81-42-995-3132)
FEATURES
  source
    Location/Qualifiers
      1..48
        /organism="Homo sapiens"
        /db_xref="taxon:9606"
        /clone="HP00302"
        /cell_line="U937"
        /tissue_type="lymphoma"
        /clone_lib="U937/pkai"
      1..42
        /replace="ttttcttttc"
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        /replace="cttttttttc"
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      1..10
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      1..10
        /replace="cttttttttc"
      1..10
        /replace="ctttcttttc"
      43..>48
        /codon_start=1
        /product="ribosomal protein L24"
        /protein_id="BAA05766.1"
        /db_xref="GI:4433237"
        /translation="MK"
BASE COUNT
ORIGIN
  5 a 13 c 10 g 20 t

alignment_scores:
  Quality: 40.00      Length: 15
  Ratio: 2.857       Gaps: 0
Percent Similarity: 93.333 Percent Identity: 46.667

alignment_block:
  US-09-836-410A-1 x HUMRPY59/rev ..
  Align seg 1/1 to reverse of: HUMRPY59 from: 1 to: 48

578 ValAsnGlyAspSerAlaGluThrGluGluLeuAlaAsnGlu 592
48 CITCATGGCGACAGCTCCACGGAACACAAAGATGCCGAAGAA 4
seq_name: gb_pat:AX147194

seq_documentation_block:
  LOCUS AX147194 50 bp DNA linear PAT 08-JUN-2001
  DEFINITION Sequence 32 from Patent WO0136682.
  ACCESSION AX147194
  VERSION AX147194.1 GI:14346365
  KEYWORDS
  SOURCE
    synthetic construct.
```


36 CCGG 39

ORGANISM synthetic construct
artificial sequence.
REFERENCE 1 (bases 1 to 50)
AUTHORS Chenchik,A., Munishkin,A. and Simonenko,P.
TITLE Long oligonucleotide arrays
JOURNAL Patent: WO 0136682-A 32 25-MAY-2001;
Clontech Laboratories Inc. (US)
FEATURES
source
1..50
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="synthetic oligonucleotide"
BASE COUNT 17 a 6 c 16 g 11 t
ORIGIN
alignment_scores:
Quality: 40.00 Length: 9
Ratio: 4.44 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 66.667
alignment_block:
US-09-836-410A-1 x AX147194/rev ..
Align seg 1/1 to reverse of: AX147194 from: 1 to: 50

428 AlaileAspSerHisProTTPLeu 436

|||||:|||||:|||||:|||||:|||||
44 GCACCTTTCTCAAGTCACCCCTTGGCTG 18

seq_name: gb_pat:AX222245

seq_documentation_block:
LOCUS AX222245 48 bp mRNA linear PAT 07-SEP-2001
DEFINITION Sequence 7687 from Patent WO0159103.
ACCESSION AX222245
VERSION AX222245.1 GI:15549969

KEYWORDS
SOURCE synthetic construct.
ORGANISM
artificial sequence.
REFERENCE 1 (bases 1 to 48)
AUTHORS Blatt,L., McSwiggen,J. and Chowrira,B.M.
TITLE Method and reagent for the modulation and diagnosis of cd20 and
nogo gene expression
JOURNAL Patent: WO 0159103-A 7687 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US) ; Blatt, Lawrence (US) ;
McSwiggen, James (US) ; Chowrira, Bharat M. (US)
FEATURES
Location/Qualifiers
source
1..48
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Nucleic Acid"

BASE COUNT 12 a 16 c 11 g 9 t
ORIGIN
alignment_scores:
Quality: 39.50 Length: 18
Ratio: 3.038 Gaps: 1
Percent Similarity: 72.222 Percent Identity: 50.000

alignment_block:
US-09-836-410A-1 x AX222245 ..

Align seg 1/1 to: AX222245 from: 1 to: 48

380 ThrProLeuGluAlaLeuLysPheLeuThrProLeuLysAsnLeuVa 396
|||||:|||||:|||||:|||||:|||||
1 ACACCACTGGAGGAA.....ACTCCCTTCAAGGACATCGT 35

396 LLys 397
|:::

seq_name: gb_pat:AX111996
seq_documentation_block:
LOCUS AX111996 30 bp DNA linear PAT 01-MAY-2001
DEFINITION Sequence 9 from Patent WO0125439.
ACCESSION AX111996
VERSION ; AX111996.1 GI:13938904
KEYWORDS ;
SOURCE synthetic construct.
ORGANISM
artificial sequence.
REFERENCE: 1 (bases 1 to 30)
AUTHORS Bonello,J.F., Rogowsky,P. and Perez,P.
TITLE Plant seed endosperm-specific promoter
JOURNAL Patent: WO 0125439-A 9 12-APR-2001;
Biogemma (FR)
FEATURES
Location/Qualifiers
source
1..30
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="SEQUENCE DESCRIPTION artificielle:oligonucleotide"
BASE COUNT 9 a 6 c 6 g 9 t
ORIGIN

alignment_scores:
Quality: 39.00 Length: 10
Ratio: 4.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
US-09-836-410A-1 x AX111996 ..

Align seg 1/1 to: AX111996 from: 1 to: 30

230 GlyGluAlaLeuLysLysCysHisGluIle 239
|||||:|||||:|||||:|||||:|||||
1 GGGGAAGCTTTACATCTTGGCATAACATA 30

seq_name: gb_pat:AR123920

seq_documentation_block:
LOCUS AR123920 47 bp DNA linear PAT 16-MAY-2001
DEFINITION Sequence 4 from patent US 6171823.
ACCESSION AR123920
VERSION AR123920.1 GI:14109281

KEYWORDS
SOURCE Unknown.
ORGANISM
Unclassified.
REFERENCE 1 (bases 1 to 47)
AUTHORS Woldike,H.Fabricius, and Hastrup,S.
TITLE Process for producing extracellular proteins in bacteria
JOURNAL Patent: US 6171823-A 4 09-JAN-2001;
FEATURES
Location/Qualifiers
source
1..47
/organism="unknown"

BASE COUNT 6 a 18 c 12 g 11 t
ORIGIN

alignment_scores:
Quality: 39.00 Length: 9
Ratio: 4.333 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 66.667

alignment_block:
US-09-836-410A-1 x AR123920/rev ..

Align seg 1/1 to reverse of: AR123920 from: 1 to: 47

355 LysLysAspAspAspGluGluLeu 363
:::|||||
27 GAGAGGACGCGATGATAAGAGGTC 1

seq_name: gb_pat:129948

seq_documentation_block:

LOCUS I29948 50 bp DNA linear PAT 06-FEB-1997

DEFINITION Sequence 12 from patent US 5578478.

ACCESSION I29948

VERSION I29948.1 GI:1820739

KEYWORDS

SOURCE

ORGANISM

REFERENCE

1 (bases 1 to 50)

Unclassified.

AUTHORS Rambosek, J., Piddington, C.S., Kovacevich, B.R., Young, K.D. and

Denome, S.A.

TITLE Recombinant DNA encoding a desulfurization biocatalyst

JOURNAL Patent: US 5578478-A 12 26-NOV-1996;

FEATURES Location/Qualifiers

source

1..50

/organism="unknown"

BASE COUNT 17 a 13 c 12 g 8 t

ORIGIN

alignment_scores:

Quality: 39.00

Ratio: 4.875

Length: 9

Gaps: 0

Percent Similarity: 88.889

Percent Identity: 66.667

alignment_block:

US-09-836-410A-1 x I29948/rev ..

Align seg 1/1 to reverse of: I29948 from: 1 to: 50

553 AlasercysHisLysLeuPheProTyr 561

|||||

43 GCGGCTGTCTCATGCTCTGTTCTCAT 17

seq_name: gb_pat:AX223522

seq_documentation_block:

LOCUS AX223522 48 bp mRNA linear PAT 07-SEP-2001

DEFINITION Sequence 8964 from Patent WO0159103.

ACCESSION AX223522

VERSION AX223522.1 GI:15551246

KEYWORDS

SOURCE

ORGANISM

synthetic construct.

synthetic construct

artificial sequence.

REFERENCE

1 (bases 1 to 48)

AUTHORS Blatt, L., McSwiggen, J. and Chowrira, B.M.

TITLE Method and reagent for the modulation and diagnosis of cd20 and

nogo gene expression

JOURNAL Patent: WO 0159103-A 8964 16-AUG-2001;

RTBOZYME PHARMACEUTICALS, INC. (US); Blatt, Lawrence (US);

MCSWIGGEN, James (US); Chowrira, Bharat M. (US)

FEATURES Location/Qualifiers

source

1..48

/organism="synthetic construct"

/db_xref="taxon:32630"

/note="Nucleic Acid"

BASE COUNT 13 a 13 c 12 g 10 t

ORIGIN

alignment_scores:

Quality: 38.50

Ratio: 2.406

Length: 21

Gaps: 1

Percent Similarity: 76.190

Percent Identity: 38.095

alignment_block:

US-09-836-410A-1 x AX223522 ..

Align seg 1/1 to: AX223522 from: 1 to: 48

380 ThrProLeuGluAlaIleLysPheLeuThrProLeuLysAsnLeuVa 396

:::|||||

1 AGCCCATGTGGAGGAA.....ACTCCCTTCAAGGACATCGT 35

396 lLysAsnLysIle 400

|||||

36 CCGGGATGATCTG 48

seq_name: gb_pat:AX229410

seq_documentation_block:

LOCUS AX229410 48 bp mRNA linear PAT 10-SEP-2001

DEFINITION Sequence 2782 from Patent WO0157206.

ACCESSION AX229410

VERSION AX229410.1 GI:15558551

KEYWORDS

SOURCE

ORGANISM

synthetic construct.

synthetic construct

artificial sequence.

REFERENCE

1 (bases 1 to 48)

AUTHORS Fattaey, A.R., Jarvis, T., Mcswiggen, J., Booher, R.N. and Holman, P.S.

TITLE Method and reagent for the inhibition of checkpoint kinase-1 (chk

1) enzyme

JOURNAL Patent: WO 0157206-A 2782 09-AUG-2001;

RIBOZYME PHARMACEUTICALS, INC. (US); Fattaey, Ali R. (US)

FEATURES Location/Qualifiers

source

1..48

/organism="synthetic construct"

/db_xref="taxon:32630"

BASE COUNT 11 a 14 c 14 g 9 t

ORIGIN

alignment_scores:

Quality: 38.50

Ratio: 2.567

Length: 21

Gaps: 1

Percent Similarity: 71.429

Percent Identity: 38.095

alignment_block:

US-09-836-410A-1 x AX229410 ..

Align seg 1/1 to: AX229410 from: 1 to: 48

380 ThrProLeuGluAlaIleLysPheLeuThrProLeuLysAsnLeuVa 396

|||||

1 ACCCTGCGGAGGAA.....ACTCCCTTCAAGGACATCGT 35

396 lLysAsnLysIle 400

|||||

36 CCGGGATGATGTTG 48

us-09-836-410a-1.p2n15to50.rge

Mon Jul 22 09:40:55 2002


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alignment_scores:
  Quality: 44.00      Length: 16
  Ratio: 3.385        Gaps: 0
  Percent Similarity: 81.250  Percent Identity: 50.000

alignment_block:
  US-09-836-410A-1 x AAF57073      ..
  Align seg 1/1 to: AAF57073      from: 1 to: 49

359 AspAspGluIleGlyGlyProIysGluLeuIleProGluLys 374
|||||:|||||:|||||  |||  |||||:|||||:|||||
1 GACGACGACAAAGATGGGTGCCCAATTGGTGAGATTATACCAAGAAAA 48

alignment_scores:
  Quality: 4.778      Length: 0
  Ratio: 90.000       Gaps: 0
  Percent Similarity: 80.000  Percent Identity: 80.000

alignment_block:
  US-09-836-410A-1 x AAT47976      ..
  Align seg 1/1 to: AAT47976      from: 1 to: 48

356 LysAspAspAspGluIleGlyGly 365
|||||:|||||:|||||  |||||
13 AAGGACGACGATGACAAGATCATCGGGGGC 42

seq_name: /SIDSL/gcgdata/hold-geneseq/geneseqn-emb1/NA2000.DAT:AAA08530

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seq_documentation_block:

ID	AAA08530	standard; DNA; 50 BP.
XX		
AC	AAA08530;	
XX		
DT	19-JUL-2000	(first entry)
XX		
DE	Oligonucleotide encoding C-terminal portion of the alpha factor signal	
DE	sequence and mutated trypsinogen leader sequence.	
XX		
KW	C-terminal alpha factor signal sequence; trypsinogen; leader sequence;	
KW	analogue; mutated bovine trypsinogen; recombinant protein production;	
KW	inactive zymogen; ss.	
XX		
OS	Synthetic.	
XX		
XX		
PN	WO200017332-A1.	
XX		
PD	30-MAR-2000.	
XX		
PF	15-SEP-1999;	99WO-US21047.
XX		
PR	21-SEP-1998;	98US-0101213.
XX		
PA	(ELIL) LILLY & CO ELI.	
XX		
PI	Hanqueler JM, Hershberger CL, Desplancq D, Larson JL, Rosteck PR;	
PI		
DR	WPI; 2000-283565/24.	
XX		
PT	New trypsinogen analog useful for the production of recombinant trypsin	
PT	has a modified leader sequence not cleavable by trypsin or trypsin-like	
PT	enzymes	
XX		
PS	Example 1; Page 29; 56pp; English.	
XX		
CC	Trypsinogen was fused directly to the C-terminus of the alpha factor	
CC	without a Glu-Ala-Glu-Ala linker peptide. Oligonucleotides AAA08529-30	
CC	encoding a C-terminal portion of the alpha factor signal sequence and	
CC	the Val(Asp)5 leader sequence were synthesized. The wild type bovine	
CC	trypsinogen was mutated to destroy the trypsin cleavage site. The lys	
CC	residue present in the leader sequence of the native bovine trypsinogen	
CC	protein was mutated to an Asp residue. The specification claims an	
CC	isolated trypsinogen analogue comprising a protein having trypsin	
CC	activity and a leader sequence having at least two amino acids which	
CC	are not Lys or Arg. A recombinantly produced trypsin (AA91926) is also	
CC	claimed. The trypsin derived from the recombinant trypsinogen is useful	
CC	for the characterization of other proteins, and in the manufacture of	
CC	other recombinant bioproducts, for example to cleave leader sequences	
CC	from small recombinant proteins expressed initially as fusion proteins.	
CC	The present method provides for expression of an inactive zymogen form	
CC	that is soluble and properly folded yet is not activated until after	
CC	purification from fermentation broth or cell extracts. This is	
CC	accomplished through the expression of a single chain trypsinogen	
CC	analogue where the leader sequence is modified such that it lacks a	
CC	trypsin-like enzyme cleavage site. Specifically the trypsinogen	
CC	analogues of the present invention lack a lysine or arginine in the	
CC	N-terminal leader sequence of the protein to prevent auto-activation or	
CC	activation by endogenous host cell enzymes.	
XX		
SQ	Sequence 50 BP; 12 A; 17 C; 6 G; 15 T; 0 other;	

```

354 LysLysLysAspAspGluGluLeuGlyGly 365
||||| ||||| ||||| ::|||
49 AAAGATCGACGATGATGACGATATCGTTGGAGT 14

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-emb1/NA2001A.DAT-AAI75479

seq_documentation_block:
ID_ AAI75479 standard; DNA; 50 BP.
XX AC
XX AC AAI75479;
XX XX
XX DT 09-NOV-2001 (first entry)
XX DE
XX DE Human silent SNP containing nucleic acid SEQ:2420.
XX KW Human; single nucleotide polymorphism; SNP; genome; gene therapy;
XX KW protein therapy; vaccine; probe; diagnostic assay; detection;
XX KW quantitation; restorative therapy; polymorphic; ds.
XX OS Homo sapiens.
XX XX
XX PN WO200140521-A2.
XX XX
XX PD 07-JUN-2001.
XX PF 30-NOV-2000; 2000WO-US32758.
XX PR 30-NOV-1999; 99US-0168138.
XX PR 29-NOV-2000; 2000US-0726173.
XX PA (CURA-) CURAGEN CORP.
XX PI Shimkets RA, Leach M;
XX DR WPI; 2001-356160/37.
XX PT Polymorphic nucleic acid sequences, useful in genetic testing and
XX PT therapy
XX PS Claim 1; Page 792; 2653pp; English.
XX CC AAI73060 to AAI79867 represent isolated human polymorphic polynucleotide
XX CC sequences (I), which contain single nucleotide polymorphisms (SNPs).
XX CC AAC53114 to AAM53329 represent peptides related to human polymorphic
XX CC polynucleotide sequences. The sequences can be used in gene and protein
XX CC therapy, and in vaccine production. (I) and the polypeptides encoded by
XX CC them may be used in the prevention, diagnosis and treatment of diseases
XX CC associated with inappropriate expression of polymorphic polypeptides.
XX CC For example, (I) may be used to treat disorders by rectifying mutations
XX CC or deletions in a patient's genome that affect the activity of
XX CC polypeptides by expressing inactive proteins or to supplement the
XX CC patients own production of polypeptide. Additionally, (I) and its
XX CC complementary sequences may also be used as DNA probes in diagnostic
XX CC assays to detect and quantify the presence of similar nucleic acids
XX CC in samples, and therefore which patients may be in need of restorative
XX CC therapy. The polypeptides encoded by (I) may be used as antigens in the
XX CC production of antibodies specific for polymorphic polypeptides. The
XX CC antibodies may also be used to down regulate expression and activity.
XX CC The antibodies may also be used as diagnostic agents for detecting the
XX CC presence of polymorphic polypeptides in samples.
XX SQ Sequence 50 BP; 10 A; 17 C; 14 G; 9 T; 0 other;

alignment_scores:
Quality: 43.00 Length: 9
Ratio: 4.778 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:
US-09-836-410A-1 x AAI75479/rev ..
Align seg 1/1 to reverse of: AAI75479 from: 1 to: 50

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alignment_block:
US-09-836-410A-1 x AAL34450
Align seg 1/1 to: AAL34450 from: 1 to: 50
6 LysIleTyrGluGluAlaTyrThrLysTyrProArgGlyLeuValProAr 22
|||||:|||||:|||||:|||||
3. AAATATATATGAGAAAGCTGGCAAC..... 26
22 glysLeuProLeuAsnPheLeuSerGlyGlu 32
|||||:|||||:|||||:|||||
27CCAGTGAACCTCTCGCGGAGAG 50

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-emb1/NA1999.DAT: AAX88379

seq_documentation_block:
ID AAX88379 standard: DNA; 45 BP.

XX AAX88379;
AC AAX88379;
DT 30-SEP-1999 (first entry)
XX Antibody lambda light chain variable region PCR primer Rjambda0-B.
DE Haematopoietic; growth factor; PCR primer; mimetic; cell survival;
KW cell-proliferation; cell differentiation; cell activation; agonist;
KW growth factor inhibitor; nervous system cell; endodermal cell; therapy;
KW totipotent cell; embryonic stem cell; gene therapy; protection; allergy;
KW diagnostic; neutropenia; leukemia; aplastic anaemia; thrombocytopenia;
KW cancer cell; bone marrow transplant; myeloproliferative disease;
KW antibody; lambda light chain; variable region; ss.
XX Synthetic.
OS WO9938008-Al.
XX 29-JUL-1999.
PN 22-JAN-1999; 99WO-US01331.
PF 23-JAN-1998; 98US-0072253.
PR (PROL-) PROLIFARON INC.
XX Bowdish KS;
XX WPI; 1999-458732/38.

Identification of agonist or inhibitory antibodies to receptors that control cellular processes, used to modulate, e.g. proliferation of hematopoietic cells

Example 1; Page 115; 123pp; English.

This invention describes a novel method for identifying agonist or inhibitory antibodies (Ab) to receptors (R) involved in cell survival, proliferation, differentiation, or activations. The method is used to identify Ab that are growth factor mimetics and inhibitors and can regulate growth, differentiation, survival and activity of many different cell types, particularly hematopoietic cells (at various stages, of any lineage), but also nervous system cells, endodermal cells or totipotent (embryonic stem) cells. Ab are used (i) directly as therapeutic agents, e.g. for amplifying particular cell types, for ex vivo proliferation or differentiation of cells for use in gene therapy, to protect normal cells against chemotherapeutic agents, and to treat conditions such as allergy, (ii) as diagnostic/research reagents, e.g. for cell identification and sorting, (iii) to clone receptors and native factors that they mimic (also potential therapeutic agents). Typical therapeutic applications are in cases of neutropenia (of any etiology) or aplastic anemia, bone marrow transplants, myeloproliferative diseases (e.g. leukemia, thrombocytopenia or allergy) and inhibitory Ab optionally coupled to a toxin, are used to kill cancer cells. The method is not subject to the usual limitations of monoclonal antibody technology

17 ArgGlyLeuValProArgLysLeuPro 25
|||||:|||||:|||||:|||||
43 AGAGGGCTGGTCCACGTCAGCTGCG 17
seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-emb1/NA2001A.DAT: AAL34450
seq_documentation_block:
ID AAL34450 standard: DNA; 50 BP.
XX AAL34450;
XX 24-JAN-2002 (first entry)
DE Human SNP oligonucleotide #7658.
XX Immunosuppressive; immunostimulatory; antiinflammatory; cytostatic;
KW neuroprotective; antimicrobial; gene therapy; vaccine; amylase; cancer;
KW amyloid protein; angiotensin; apoptosis related protein; cadherin;
KW cyclin; polymerase; oncogene; histone; kinase; colony stimulating factor;
KW complement related protein; cytochrome; kinesin; cytokine; interferon;
KW interleukin; G-protein coupled receptor; thioesterase; inflammation;
KW multifactorial disease; autoimmune disease; infection;
KW nervous system disease; ss.
XX Homo sapiens.
OS WO200147944-A2.
XX 05-JUL-2001.
XX 28-DEC-2000; 2000WO-US35498.
XX 28-DEC-1999; 99US-0173419.
PR 27-DEC-2000; 2000US-0173419.
XX (CURA-) CURAGEN CORP.
XX Shimkets RA, Leach M;
PI WPI; 2001-465210/50.
XX Polymorphic nucleic acids encoding e.g. amylases, cyclins, polymerases,
XX oncogenes and histones, useful for diagnosing and treating, e.g.
XX cancer, autoimmune diseases and infections -
XX Claim 1; Page 3600; 4143pp; English.

The present invention relates to oligonucleotides encoding polymorphic variants of proteins related to amylases, amyloid proteins, angiotensin, apoptosis related proteins, cadherin, cyclin, polymerase, oncogenes, histones, kinases, colony stimulating factors, complement related proteins, cytochromes, kinesins, cytokines, interferons, interleukins, G-protein coupled receptors and thioesterases. The present sequence is one such oligonucleotide. The oligonucleotides and the peptides encoded by them may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate expression of the proteins listed above. Disorders that may be prevented, diagnosed and/or treated include multifactorial diseases with a genetic component, such as autoimmune diseases (e.g. rheumatoid arthritis, multiple sclerosis, diabetes, cancer systemic lupus erythematosus and Grave's disease), inflammation, cancer (e.g. cancers of the bladder, brain, breast, colon and kidney, leukaemia), diseases of the nervous system and an infection of pathogenic organisms.

Sequence 50 BP; 18 A; 11 C; 13 G; 8 T; 0 other;

alignment_scores:
Quality: 42.50 Length: 27
Ratio: 2.833 Gaps: 1
Percent Similarity: 55.556 Percent Identity: 37.037

PS Claim 1; Page 551; 2745pp; English.

XX AAZ65654 to AAZ69578 represent human biallelic markers from the present
 CC invention, which contain a polymorphic base at position 24 of their
 CC nucleotide sequences. AAZ69579 to AAZ77440 represent amplification
 CC primers for the biallelic markers. The biallelic markers of the
 CC invention have a variety of uses: they can be used for high density
 CC mapping of the human genome, and in complex association studies and
 CC haplotyping studies which are useful in determining the genetic basis
 CC for disease states. Compositions and methods of the invention can also
 CC be useful for the identification of the targets for the development of
 CC pharmaceutical agents and diagnostic methods, as well as the
 CC characterisation of the differential efficacious responses to and side
 CC effects from pharmaceutical agents acting on a disease as well as other
 CC treatment.

CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
 CC and 3367, are not actually given a sequence in the Sequence Listing
 CC from the present invention.

XX
 SQ Sequence 47 BP; 19 A; 5 C; 10 G; 13 T; 0 other;

alignment_scores:
 Quality: 42.00 Length: 10
 Ratio: 5.250 Gaps: 0
 Percent Similarity: 80.000 Percent Identity: 70.000

alignment_block:
 US-09-836-410A-1 x AAZ67157 ..

Align seg 1/1 to: AAZ67157 from: 1 to: 47

6 LysileTyrGluGluAlaTrpThrLysTyr 15
 17 AAATATATATGACAGACAGACTGGCCAAATAT 46

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-embl/NA2001A.DAT:AAH75873

seq_documentation_block:
 ID AAH75873 standard; DNA; 41 BP.

XX
 AC AAH75873;
 DT 26-OCT-2001 (first entry)
 XX
 DE Human reverse transcriptase 13 coding sequence probe #1.
 XX
 KW Human; reverse transcriptase 13; cytostatic; virucide; immunomodulatory;
 KW antiinflammatory; haemostatic; gene therapy; malignant tumour;
 KW haemopathy; HIV infection; immunological disease; inflammation;
 KW developmental disorder; probe; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200164893-A1.
 XX
 PD 07-SEP-2001.
 XX
 PF 26-FEB-2001; 2001WO-CN00280.
 XX
 PR 02-MAR-2000; 2000CN-0111806.
 XX
 PA (BIOW-) BIOWINDOW GENE DEV INC SHANGHAI.
 XX
 PI Mao Y, Xie Y;
 XX
 DR WPI; 2001-550183/61.
 XX
 CC New human reverse transcriptase 13 for diagnosing and treating
 PT developmental disorders, malignant tumor, hemopathy, human
 PT immunodeficiency virus infection, immunological diseases and
 PT inflammations

PS Example 7; Page 15; 34pp; Chinese.

XX The present invention relates to human reverse transcriptase 13 and its
 CC coding sequence (see AAH75868 and AAG66428). The reverse transcriptase
 CC and its coding sequence are useful in the diagnosis and treatment of
 CC malignant tumour, haemopathy, HIV infection, immunological diseases,
 CC various inflammations and developmental disorders. The present sequence
 CC is a probe, which was used in an example from the present invention.

XX
 SQ Sequence 41 BP; 18 A; 7 C; 6 G; 10 T; 0 other;

alignment_scores:
 Quality: 41.00 Length: 14
 Ratio: 3.417 Gaps: 1
 Percent Similarity: 85.714 Percent Identity: 42.857

alignment_block:
 US-09-836-410A-1 x AAH75873/rev ..

Align seg 1/1 to reverse of: AAH75873 from: 1 to: 41

435 TrpLeuHisGluCysMetIleArgLeuPheHisSerValCys 448
 39 TGGCTGCATAAATGTCTT.....CTTTGAGAAATATCTGT 4

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-embl/NA1999.DAT:AAK05908

seq_documentation_block:
 ID AAK05908 standard; DNA; 47 BP.

XX
 AC AAK05908;
 DT 07-MAY-1999 (first entry)
 XX
 DE Oligonucleotide probe meca945-29A18P.
 XX
 KW Hybridization; RNase H; scissile linkage; nucleic acid detection;
 KW gene detection; polyamine; probe; DNA/RNA hybrid; ss.
 XX
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT misc_RNA 14..17
 FT /*tag= a
 XX
 PN WO9901570-A2.
 XX
 PD 14-JAN-1999.
 XX
 PF 03-JUL-1998; 98WO-CA00631.
 XX
 PR 22-JUN-1998; 98US-0090273.
 PR 03-JUL-1997; 97US-0051827.
 PR 18-MAY-1998; 98US-0086021.
 XX
 PA (IDBI-) ID BIOMEDICAL CORP.
 XX
 PI Bryan RN, Cloney LP, Farnworth BA, Marostenmaki AJ;
 XX
 DR WPI; 1999-106070/09.
 XX
 CC Increasing the hybridization rate between two nucleic acids - using
 PT ribonuclease H (RNase H) and/or a polyamine, useful for detecting
 PT nucleic acids of interest in a sample
 XX
 PS Example 1; Page 15; 45pp; English.
 XX
 CC The invention relates to methods of increasing the hybridization rate
 CC between two nucleic acids. One method comprises construction of two
 CC nucleic acids and a polyamine, and hybridizing both nucleic acids
 CC together, under suitable conditions. Also provided is a similar method
 CC involving two nucleic acids with RNase H, where both nucleic acids do

CC not contain a scissile linkage, and if one nucleic acid is DNA, then the
 CC other is RNA. The methods are useful for (in)directly detecting nucleic
 CC acids of interest in a sample. They may be applied to short and long
 CC nucleic acids. The acceleration of rate reactions in gene detection
 CC assays, using RNase H and/or polyamines (under conditions of low salt
 CC concentration), produces a lower background and, therefore, a greater
 CC signal to noise ratio. Sequences AAX05903-914 represent oligonucleotide
 CC probes used in the course of the invention.
 XX
 SQ Sequence 47 BP; 37 A; 1 C; 7 G; 2 T; 0 other;

alignment_scores:
 Quality: 41.00 Length: 15
 Ratio: 3.727 Gaps: 0
 Percent Similarity: 73.333 Percent Identity: 60.000

alignment_block:

US-09-836-410A-1 x AAX05908 ..

Align seg 1/1 to: AAX05908 from: 1 to: 47

342 AsnAlaGluLysGluLysProGlnArgAsnProLysLysLys 356
 ||| |||||
 1 AATAGAGAAAAAGAAAAAGATGCGCAAGAAAAA 45

seq_name: /SIDSI/gcgdata/hold-geneseq/geneseq-n-emb1/NA1999.DAT: AAX15480

seq_documentation_block:

ID AAX15480 standard; DNA; 47 BP.

XX AC AAX15480;

XX DT 07-MAY-1999 (first entry)

XX DE Probe mecA945-29A18P for detecting an antibiotic resistant mecA gene.

XX KW Antibiotic resistant mecA gene; transmission; treatment;

XX KW methicillin resistant; Staphylococcus; DNA/RNA hybrid; probe; ss.

XX OS Synthetic.

XX OS Staphylococcus sp.

XX FH Key Location/Qualifiers

XX FT misc_RNA 14..17

XX FT /*tag= a

XX PN WO9901572-A2.

XX PD 14-JAN-1999.

XX PF 03-JUL-1998; 98WO-CA00633.

XX PR 22-JUN-1998; 98US-0090276.

XX PR 03-JUL-1997; 97US-0051643.

XX PR 18-MAY-1998; 98US-0086020.

XX PA (IDBI-) ID BIOMEDICAL CORP.

XX PI Bekkaoui F, Cloney LP;

XX DR WPI; 1999-106072/09.

XX Method for determining the presence of an antibiotic resistant mecA
 PT gene in a sample - using a scissile link containing nucleic acid
 PT probe for antibiotic resistant mecA gene

XX Example 1; Page 16; 59pp; English.

XX The present sequence represents a probe used for determining the
 CC presence of an antibiotic resistant mecA gene in a biological sample.
 CC The method provides a means for the rapid detection, for both
 CC the prevention of transmission and treatment of, methicillin resistant

CC Staphylococcus species.

XX SQ Sequence 47 BP; 37 A; 1 C; 7 G; 2 T; 0 other;

alignment_scores:

Quality: 41.00 Length: 15
 Ratio: 3.727 Gaps: 0
 Percent Similarity: 73.333 Percent Identity: 60.000

alignment_block:

US-09-836-410A-1 x AAX15480 ..

Align seg 1/1 to: AAX15480 from: 1 to: 47

342 AsnAlaGluLysGluLysProGlnArgAsnProLysLysLys 356
 ||| |||||
 1 AATAGAGAAAAAGAAAAAGATGCGCAAGAAAAA 45

seq_name: /SIDSI/gcgdata/hold-geneseq/geneseq-n-emb1/NA2000.DAT: AAX67285

seq_documentation_block:

ID AAX67285 standard; DNA; 47 BP.

XX AC AAX67285;

XX DT 10-SEP-2001 (first entry)

XX DE Human map-related biallelic marker SEQ ID NO:1632.

XX KW Human genome; biallelic marker; high density disequilibrium map;
 KW genomic map; haplotype; phenotype; polymorphic base; genotyping;
 KW haplotyping; hybridisation; identification; characterisation;
 KW diagnosis; single nucleotide polymorphism; SNP; ds.

XX OS Homo.sapiens.

XX FH Key Location/Qualifiers

XX FT variation replace(24,C)

XX FT /*tag= a

XX FT /standard_name= "single nucleotide polymorphism"

XX PN WO9954500-A2.

XX PD 28-OCT-1999.

XX PF 21-APR-1999; 99WO-IB00822.

XX PR 21-APR-1998; 98US-0082614.

XX PR 23-NOV-1998; 98US-0109732.

XX PA (GEST) GENSET.

XX PI Cohen D, Blumenfeld M, Chumakov I;

XX DR WPI; 2000-013267/01.

XX Novel biallelic markers used to construct a high density disequilibrium
 PT map of the human genome -

XX Claim 1; Page 578; 2745pp; English.

XX AZ65654 to AZ69578 represent human biallelic markers from the present
 CC invention, which contain a polymorphic base at position 24 of their
 CC nucleotide sequences. AZ69579 to AZ77440 represent amplification
 CC primers for the biallelic markers. The biallelic markers of the
 CC invention have a variety of uses: they can be used for high density
 CC mapping of the human genome, and in complex association studies and
 CC haplotyping studies which are useful in determining the genetic basis
 CC for disease states. Compositions and methods of the invention can also
 CC be useful for the identification of the targets for the development of
 CC pharmaceutical agents and diagnostic methods, as well as the
 CC characterisation of the differential efficacious responses to and side

CC effects from pharmaceutical agents acting on a disease as well as other
CC treatment.
CC N.B. The SEQ ID NOS 2852, 2913, 2974, 3035, 3096, 3157, 3227, 3297
CC and 3367, are not actually given a sequence in the Sequence Listing
CC from the present invention.

XX Sequence 47 BP; 20 A; 6 C; 10 G; 11 T; 0 other;

alignment_scores:
Quality: 41.00 Length: 15
Ratio: 4.556 Gaps: 0
Percent Similarity: 60.000 Percent Identity: 46.667

alignment_block:
US-09-836-410A-1 x AAZ67285/rev ..

Align seg 1/1 to reverse of: AAZ67285 from: 1 to: 47

97 ProProThrThrLeuLeuTrpValGlnTyrTyrLeuAlaGlnHis 111
||| ||| ::|||::|||::||| ||| ||| |||
47 CCGACACCTTATGCTGTGGATTTTACATTTTACATTTTACAC 3

seq_name: /STD1/gcgdata/hold-geneseq/geneseq-emb1/NA1989.DAT:AA92016

seq_documentation_block:
ID AAN92016 standard; DNA; 50 BP.

XX
AC AAN92016;

DT 17-APR-1990 (first entry)

XX Sequence probe complementary to Neisseria gonorrhoeae genomic sequence
DE SSJK1 combined with the LIA2C amplifier sequence.

XX Neisseria gonorrhoeae genomic sequence SSJK1; LIA2C amplifier sequence;
KW file 'rcjk'; jkl.probes15(50).

XX Neisseria gonorrhoeae.

XX Key Location/Qualifiers
FH 1..30
FT misc_feature /*tag= a
FT /*sequence probe"
FT 31..50
FT misc_feature /*tag= b
FT /*LIA2C amplifier sequence"

XX W08903891-A.

XX 05-MAY-1989.

XX 14-OCT-1988; 88WO-US03644.

XX 30-SEP-1988; 88US-0252638, US-109282.

XX (CHIR-) CHIRON CORP.

XX Urdea MS, Warner B, Running JA, Kolberg JA, Clyne JM;
PI Sanchez-Pescador R;

XX WPI; 1989-150787/20.

XX Nucleic acid multimer for hybridisation assays
PT - having single-stranded oligo-nucleotide units
PT capable of binding specifically to sequences of interest.

XX Fig 14; : 112pp; English.

XX The sequence probe (tag a) is complementary to N. gonorrhoeae genomic
CC sequence SSJK1 from the file 'rcjk'. It is used to assay crude cellular
CC lysates and genomic DNA from different bacteria.. It is called
CC jkl.probes15(50).

XX Sequence 50 BP; 11 A; 12 C; 13 G; 14 T; 0 other;

alignment_scores:
Quality: 41.00 Length: 13
Ratio: 4.100 Gaps: 0
Percent Similarity: 76.923 Percent Identity: 53.846

alignment_block:
US-09-836-410A-1 x AAN92016/rev ..

Align seg 1/1 to reverse of: AAN92016 from: 1 to: 50

519 GlySerLeuThrAsnArgAsnLeuGlnThrCysMetGlu 531
||| ::|||::|||::||| ||| ||| |||
44 GGTCCTATGCTTAATCAGAATCTGCATATCTGCATGGAG 6

seq_name: /STD1/gcgdata/hold-geneseq/geneseq-emb1/NA2001A.DAT:AA134455

seq_documentation_block:
ID AAL34455 standard; DNA; 50 BP.

XX
AC AAL34455;

DT 24-JAN-2002 (first entry)

XX Human SNP oligonucleotide #7663.

XX Immunosuppressive; immunostimulatory; antiinflammatory; cytostatic;
KW neuroprotective; antimicrobial; gene therapy; vaccine; amylase; cancer;
KW amyloid protein; angiotensin; apoptosis related protein; cadherin;
KW cyclin; polymerase; oncogene; histone; kinase; colony stimulating factor;
KW complement related protein; cytochrome; kinesin; cytokine; interferon;
KW interleukin; G-protein coupled receptor; thioesterase; inflammation;
KW multifactorial disease; autoimmune disease; infection;
KW nervous system disease; ss.

XX Homo sapiens.

XX W0200147944-A2.

XX 05-JUL-2001.

XX 28-DEC-2000; 2000WO-US35498.

XX 28-DEC-1999; 99US-0173419.

XX 27-DEC-2000; 2000US-0173419.

XX (CURA-) CURAGEN CORP.

XX Shimkets RA, Leach M;

XX WPI; 2001-465210/50.

XX Polymorphic nucleic acids encoding e.g. amylases, cyclins, polymerases,
PT oncogenes and histones, useful for diagnosing and treating, e.g.
PT cancer, autoimmune diseases and infections -

XX Claim 1; Page 3601; 4143pp; English.

XX The present invention relates to oligonucleotides encoding polymorphic
CC variants of proteins related to amylases, amyloid proteins, angiotensin,
CC apoptosis related proteins, cadherin, cyclin, polymerase, oncogenes,
CC histones, kinases, colony stimulating factors, complement related
CC proteins, cytochromes, kinesins, cytokines, interferons, interleukins,
CC G-protein coupled receptors and thioesterases. The present sequence is
CC one such oligonucleotide. The oligonucleotides and the peptides encoded
CC by them may be used in the prevention, diagnosis and treatment of
CC diseases associated with inappropriate expression of the proteins listed
CC above. Disorders that may be prevented, diagnosed and/or treated include
CC multifactorial diseases with a genetic component, such as autoimmune
CC diseases (e.g. rheumatoid arthritis, multiple sclerosis, diabetes,

CC systemic lupus erythematosus and Grave's disease), inflammation, cancer
 CC (e.g. cancers of the bladder, brain, breast, colon and kidney,
 CC leukaemia), diseases of the nervous system and an infection of pathogenic
 CC organisms.
 XX

SQ Sequence 50 BP; 10 A; 13 C; 22 G; 5 T; 0 other;

alignment_scores:

Quality:	41.00	Length:	12
Ratio:	4.100	Gaps:	0
Percent Smilarity:	83.333	Percent Identity:	66.667

alignment_block:

US-09-836-410A-1 x AAL34455 ..

Align seg 1/1 to: AAL34455 from: 1 to: 50

364 GlyGlyProLysGluLeuLeuProGluLysLeu 375

||||||| |||||||:|:|:|:|:|

14 GGAGGGCCGCCAGAGGAGTGCTGCGGAGACCTA 49

OM of: US-09-836-410A-1 to: EST:* out_format : pfs

Date: Jul 20, 2002 4:12 AM

About: Results were produced by the GenCore software, version 4.5,
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Command line parameters:

-MODEL=frame+p2n.model -DEV=xlh
-O=/sgn2_1/USPRO-spool/US09836410/runat_18072002_115032_29691/app_query.fasta_1.660
-DB=EST -QFMT=fastcap -SUFFIX=p2n15to50.rst -GAPOP=12.000
-CGAPEXT=4.000 -MINMATCH=0.100 -LOOPCL=0.000 -LOOPEXT=0.000
-QGAPOP=4.500 -QGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500
-FGAPOP=6.000 -FGAPEXT=7.000 -XGAPOP=10.000 -XGAPEXT=0.500
-DELOP=6.000 -DELEXT=7.000 -START=1 -MATRIX=blossum62
-TRANS=human40.cgi -LIST=45 -DOCALIGN=200 -THR_SCORE=pct
-THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=pfs
-NORM=ext -HEAPSIZE=500 -MINLEN=15 -MAXLEN=50
-USER=US09836410 -CGN1_1_2960 -NCPU=6 -ICPU=3 -LONGLOG
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -NO_XLPXY -WAIT -THREADS=1

Search information block:

Query: US-09-836-410A-1
Query length: 593
Database: EST:*
Database sequences: 13736207
Database length: 1841457050
Search time (sec): 1532.560000

score_list:

Sequence	Strd	Orig	ZScore	EScore	Len	Documentation
gb_gss:A2774479	-	48.00	92.20	6.9e+04	33	A2774479 2M0004A05F Mouse 10kb F
gb_est1:AI657570	-	47.50	85.72	9.5e+04	40	AI657570 fc15c02.y1 Zebrafish W
gb_est1:AI223998	+	47.00	87.71	1.2e+05	46	AI223998 qx12h07.x1 NCI CGAP Lym
gb_est2:BF054800	+	45.00	83.94	2.0e+05	50	BF054800 7171g09.y1 NCI CGAP Brn
gb_est1:AA590944	+	43.00	80.92	2.9e+05	50	AA590944 vm25f02.r1 Knowles Soli
gb_est1:AI813747	+	42.00	82.13	2.5e+05	37	AI813747 wk79a02.x1 NCI CGAP Pan
gb_est2:BM393347	-	41.00	78.09	4.2e+05	49	BM393347 50071-2-9-B10.f.1 Child
gb_est2:BM395447	-	41.00	78.09	4.2e+05	49	BM395447 50072-2-9-B10.f.1 Child
gb_est1:AU104189	+	41.00	77.91	4.3e+05	50	AU104189 AU104189 Sugano Homo sa
gb_est1:AU107491	+	41.00	77.91	4.3e+05	50	AU107491 AU107491 Sugano Homo sa
gb_gss:AZ860401	-	40.50	78.10	4.2e+05	45	AZ860401 2M0166E22F Mouse 10kb F
gb_est1:AI587842	+	40.00	77.76	4.3e+05	43	AZ860401 2M0166E22F Mouse 10kb F
gb_est1:AU103783	-	40.00	77.55	4.5e+05	44	AL587842 AL587842 BP Chicken Bra
gb_est1:AU104282	-	40.00	76.40	5.2e+05	50	AU103783 AU103783 Sugano Homo sa
gb_est1:AU107535	+	40.00	76.40	5.2e+05	50	AU104282 AU104282 Sugano Homo sa
gb_est1:AA574989	-	39.50	76.20	5.3e+05	50	AZ574989 vm34a03.r1 Knowles Soli
gb_gss:AZ582769	-	39.50	75.83	5.6e+05	47	AZ582769 vm34a03.r1 Knowles Soli
gb_est1:AI500599	+	39.50	75.83	5.6e+05	49	AI500599 tp93407.x1 NCI CGAP UC4
gb_est2:BJ043594	+	39.00	75.84	5.8e+05	45	BJ043594 E043594 N18B Mochii nc
gb_est1:BI330882	+	39.00	75.45	5.8e+05	47	BI330882 602981270f1 NCI CGAP LU
gb_est1:AI627881	+	39.00	75.07	6.1e+05	49	AI627881 AL627881 XGC-gastrula S
gb_est1:AA254893	+	39.00	74.89	6.3e+05	50	AA254893 mx78d07.r1 Soares mouse
gb_est1:AA366391	+	39.00	74.89	6.3e+05	50	AA366391 w4f01a1.r1 Aspergillus
gb_est1:AU107471	+	39.00	74.89	6.3e+05	50	AU107471 AU107471 Sugano Homo sa
gb_est1:AU107473	+	39.00	74.89	6.3e+05	50	AU107473 AU107473 Sugano Homo sa
gb_est1:AU107474	+	39.00	74.89	6.3e+05	50	AU107474 AU107474 Sugano Homo sa
gb_est1:AU107475	+	39.00	74.89	6.3e+05	50	AU107475 AU107475 Sugano Homo sa
gb_est1:AU107481	+	39.00	74.89	6.3e+05	50	AU107481 AU107481 Sugano Homo sa
gb_est1:AU107483	+	39.00	74.89	6.3e+05	50	AU107483 AU107483 Sugano Homo sa
gb_est1:AU107486	+	39.00	74.89	6.3e+05	50	AU107486 AU107486 Sugano Homo sa
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gb_est1:AU107490	+	39.00	74.89	6.3e+05	50	AU107490 AU107490 Sugano Homo sa
gb_est1:AU107492	+	39.00	74.89	6.3e+05	50	AU107492 AU107492 Sugano Homo sa
gb_est1:AU107493	+	39.00	74.89	6.3e+05	50	AU107493 AU107493 Sugano Homo sa
gb_est1:AU107494	+	39.00	74.89	6.3e+05	50	AU107494 AU107494 Sugano Homo sa
gb_est1:AU107495	+	39.00	74.89	6.3e+05	50	AU107495 AU107495 Sugano Homo sa
gb_est1:AU107496	+	39.00	74.89	6.3e+05	50	AU107496 AU107496 Sugano Homo sa
gb_est1:AU107497	+	39.00	74.89	6.3e+05	50	AU107497 AU107497 Sugano Homo sa

gb_est1:AU107499 + 39.00 74.89 6.3e+05 50 i AU107499 AU107499 Sugano Homo
gb_est1:AU107500 + 39.00 74.89 6.3e+05 50 i AU107500 AU107500 Sugano Homo
gb_est1:AU107501 + 39.00 74.89 6.3e+05 50 i AU107501 AU107501 Sugano Homo
gb_est1:AU107502 + 39.00 74.89 6.3e+05 50 i AU107502 AU107502 Sugano Homo
gb_est1:AU107503 + 39.00 74.89 6.3e+05 50 i AU107503 AU107503 Sugano Homo

seq_name: gb_gss:A2774479

seq_documentation_block:

LOCUS A2774479 33 bp DNA linear GSS 16-FEB-2001
DEFINITION 2M0004A05F Mouse 10kb plasmid UUGCLM library Mus musculus genomic
clone UUGC2M0004A05 F, DNA sequence.

ACCESSION A2774479.1 GI:12899972

VERSION GSS.

KEYWORDS house mouse.

SOURCE house mouse.

ORGANISM Mus musculus

REFERENCE Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 33)

AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D., Weiss,R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

Unpublished (2000)

Contact: Robert B. Weiss

University of Utah Genome Center

University of Utah

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT

84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0004 row: A column: 05

Seq primer: CGTTGTAACACGACGCCAGT

Class: plasmid ends

High quality sequence step: 33.

Location/Qualifiers

1..33

/organism="Mus musculus"

/strain="C57BL/6J"

/db_xref="taxon:10090"

/clone="UUGC2M0004A05"

/sex="Male"

/lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"

/note="Vector: PWD42nv; Purified genomic DNA from M.

musculus C57BL/6J (male) was obtained from the Jackson

Laboratory Mouse DNA Resource

(http://www.jax.org/resources/documents/dnares/). The DNA

was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA

was blunt end-repaired with T4 DNA polymerase and T4

polynucleotide kinase. Adaptor oligonucleotides were

ligated to the blunt ends in high molar excess. The

adapted DNA was purified and size-selected for a 9.5 to

10.5 kb range using preparative agarose gel

electrophoresis. Vector DNA was prepared from a derivative

of PWD42 (g1147321141g1/AP129072.1), a copy-number

inducible derivative of plasmid R1. The vector was ligated

with adaptors complementary to the insert adaptors and

purified. The sheared, adapted mouse DNA was annealed to

adapted vector DNA, and transformed into

chemically-competent E. coli XL10-Gold (Stratagene) cells

and selected for ampicillin resistance."

0 a 0 c 6 g 27 t

BASE COUNT

ORIGIN

alignment_scores:

analysis were selected following oligonucleotide hybridization fingerprinting of arrayed clones from zebrafish late somitogenesis (26 ss), adult liver or embryonic shield stage (5.6 h) libraries. Fingerprint data were used to computationally cluster cDNAs, and a single cDNA from each cluster was chosen for sequencing. In some cases multiple members of the same cluster were sequenced to assess clustering parameters or single clones were sequenced additional times to assess quality control."

BASE COUNT 8 a 11 c 14 g 7 t
ORIGIN

alignment_scores:
Quality: 47.50 Length: 11
Ratio: 4.750 Gaps: 1
Percent Similarity: 90.909 Percent Identity: 90.909

alignment_block:
US-09-836-410A-1 x AI657570/rev ..

Align seg 1/1 to reverse of: AI657570 from: 1 to: 40

560 ProTyAlaLeuAlaPheMetProGlyTyr 570
|||||
33 CCATATGCGGTACCC...ATGCCGCTGGGTAC 4

seq_name: gb_estl:AI223998

seq_documentation_block:
LOCUS AI223998 46 bp mRNA linear EST 21-DEC-1998
DEFINITION qx12h07.x1 NCI_CGAP_Lym12 Homo sapiens cDNA clone IMAGE:2001181 3' similar to TR:Q04154 Q04154 SALIVARY PROLINE-RICH PROTEIN RPL5 PRECURSOR. ; contains element MER22 repetitive element ;, mRNA sequence.

ACCESSION AI223998
VERSION AI223998.1 GI:3806711
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 46)
REFERENCE NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP).
TITLE Tumor Gene Index
JOURNAL: Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
unknown library type
Trace considered overall poor quality
Insert Length: 1534 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 1.
Location/Qualifiers
1. .46
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2001181"
/clone_lib="NCI_CGAP_Lym12"
/tissue_type="lymphoma, follicular mixed small and large cell"
/lab_host="DH10B"
/note="Organ: lymph node; Vector: pCMV-SPOPT6; Site:1: SalI; Site:2: NotI; Cloned unidirectionally. Primer: Oligo dT. Average insert size 1.25 kb. Life Technologies catalog #: 11547-015"

BASE COUNT 19 a 17 c 10 g 0 t
ORIGIN

alignment_scores:

Quality: 48.00 Length: 10
Ratio: 4.800 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 80.000

alignment_block:
US-09-836-410A-1 x AZ774479/rev ..

Align seg 1/1 to reverse of: AZ774479 from: 1 to: 33

347 LysProGlnArgAsnProLysLysLysLys 356
|||||
32 AAACCAAAAAAACCCCAAAAAA 3

seq_name: gb_estl:AI657570

seq_documentation_block:
LOCUS AI657570 40 bp mRNA linear EST 07-JUN-2001
DEFINITION fc15c02.y1 zebrafish Washu MPIMG EST Danio rerio cDNA clone IMAGE:3721442 5' similar to TR:O13017 O13017 WINGED HELIX TRANSCRIPTIONAL FACTOR MFH-1. ;, mRNA sequence.

ACCESSION AI657570
VERSION AI657570.1 GI:4755238
KEYWORDS EST.
SOURCE zebrafish.
ORGANISM Danio rerio
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes ; Cyprinidae; Danio.
1 (bases 1 to 40)
REFERENCE Clark,M., Johnson,S.L., Lehrach,H., Lee,R., Li,F., Marra,M., Eddy S., Hillier,L., Kucaba,T., Martin,J., Beck,C., Wylie,T., Underwood K., Steptoe,M., Theising,B., Allen,M., Bowers,Y., Person,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter,E., Kohn,S., Shin,F., Jackson,Y., Cardenas,M., McCann,R., Waterston,R. and Wilson,R.
Washu Zebrafish EST Project 1998
Unpublished (1998)
Contact: Stephen L. Johnson
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: zbrfish@watson.wustl.edu
cDNA Library Preparation: Matthew Clark. cDNA Library Arrayed by: Matthew Clark. DNA Sequencing by: Washington University Genome Sequencing Center Clone Distribution: Genome Systems, St. Louis, Missouri (web address: www.genomesystems.com) (email contact: info@genomesystems.com) and Research Genetics, Huntsville, Alabama (web address: www.resgen.com) (email contact: info@resgen.com) and RessourcenzentrumPrimarDatenbank, Berlin, Germany (web address: www.rzpd.de)
Trace considered overall poor quality
Possible reversed clone: similarity on wrong strand
Seq primer: T3 ET from Amersham
High quality sequence stop: 1.
Location/Qualifiers
1. .40
/organism="Danio rerio"
/db_xref="taxon:7955"
/clone="IMAGE:3721442"
/clone_lib="Zebrafish Washu MPIMG EST"
/sex="mixed"
/tissue_type="26 somite embryos, adult livers, shield stage embryos"
/lab_host="X1-blue MRF"
/note="Vector: pSPORT1; Site:1: NotI; Site:2: SalI; 1st strand cDNA was primed with a Not I oligo(dT)15 primer [5'pGACTACTTCTAGATCGAGCGCGCCCTTTTITTTT3]; double-stranded cDNA was ligated to Sal I adaptors (BRL), digested with Not I and cloned into the Not I and Sal I sites of the pSPORT1 vector (BRL). Library was constructed by Matthew Clark (Lehrach lab; ICRF, London and Max Planck Institut fuer Molekulare Genetik, Berlin). cDNAs for EST

FEATURES
source
1. .40
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2001181"
/clone_lib="NCI_CGAP_Lym12"
/tissue_type="lymphoma, follicular mixed small and large cell"
/lab_host="DH10B"
/note="Organ: lymph node; Vector: pCMV-SPOPT6; Site:1: SalI; Site:2: NotI; Cloned unidirectionally. Primer: Oligo dT. Average insert size 1.25 kb. Life Technologies catalog #: 11547-015"

BASE COUNT 19 a 17 c 10 g 0 t
ORIGIN

alignment_scores:

FEATURES
source
1. .40
/organism="Danio rerio"
/db_xref="taxon:7955"
/clone="IMAGE:3721442"
/clone_lib="Zebrafish Washu MPIMG EST"
/sex="mixed"
/tissue_type="26 somite embryos, adult livers, shield stage embryos"
/lab_host="X1-blue MRF"
/note="Vector: pSPORT1; Site:1: NotI; Site:2: SalI; 1st strand cDNA was primed with a Not I oligo(dT)15 primer [5'pGACTACTTCTAGATCGAGCGCGCCCTTTTITTTT3]; double-stranded cDNA was ligated to Sal I adaptors (BRL), digested with Not I and cloned into the Not I and Sal I sites of the pSPORT1 vector (BRL). Library was constructed by Matthew Clark (Lehrach lab; ICRF, London and Max Planck Institut fuer Molekulare Genetik, Berlin). cDNAs for EST

FEATURES
source
1. .40
/organism="Danio rerio"
/db_xref="taxon:7955"
/clone="IMAGE:3721442"
/clone_lib="Zebrafish Washu MPIMG EST"
/sex="mixed"
/tissue_type="26 somite embryos, adult livers, shield stage embryos"
/lab_host="X1-blue MRF"
/note="Vector: pSPORT1; Site:1: NotI; Site:2: SalI; 1st strand cDNA was primed with a Not I oligo(dT)15 primer [5'pGACTACTTCTAGATCGAGCGCGCCCTTTTITTTT3]; double-stranded cDNA was ligated to Sal I adaptors (BRL), digested with Not I and cloned into the Not I and Sal I sites of the pSPORT1 vector (BRL). Library was constructed by Matthew Clark (Lehrach lab; ICRF, London and Max Planck Institut fuer Molekulare Genetik, Berlin). cDNAs for EST

FEATURES
source
1. .40
/organism="Danio rerio"
/db_xref="taxon:7955"
/clone="IMAGE:3721442"
/clone_lib="Zebrafish Washu MPIMG EST"
/sex="mixed"
/tissue_type="26 somite embryos, adult livers, shield stage embryos"
/lab_host="X1-blue MRF"
/note="Vector: pSPORT1; Site:1: NotI; Site:2: SalI; 1st strand cDNA was primed with a Not I oligo(dT)15 primer [5'pGACTACTTCTAGATCGAGCGCGCCCTTTTITTTT3]; double-stranded cDNA was ligated to Sal I adaptors (BRL), digested with Not I and cloned into the Not I and Sal I sites of the pSPORT1 vector (BRL). Library was constructed by Matthew Clark (Lehrach lab; ICRF, London and Max Planck Institut fuer Molekulare Genetik, Berlin). cDNAs for EST

Quality: 47.00 Length: 19
 Ratio: 4.273 Gaps: 1
 Percent Similarity: 57.895 Percent Identity: 52.632

alignment_block:

US-09-836-410A-1 x AI223998 ..

Align seg 1/1 to: AI223998 from: 1 to: 46

```

348 proGlnArgAsnProLysLysLysLysAspAspGluGluIleG1 364
|||||:|||||
14 CCCCCAAAACCCCAAAAAA.....GG 39
364 yGlyPro 366
|||||
40 GGGGCC 46

```

seq_name: gb_est2:BF054800

seq_documentation_block:

LOCUS BF054800 50 bp mRNA linear EST 16-OCT-2000
 DEFINITION 7171d09.y1 NCI_CGAP_Brn20 Homo sapiens cDNA clone IMAGE:3340145 5',
 mRNA sequence.

ACCESSION BF054800

VERSION BF054800.1 GI:10808696

KEYWORDS

EST.

SOURCE

human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

1 (bases 1 to 50)

NCI/NINDS-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

National Cancer Institute / National Institute of Neurological

Disorders and Stroke, Brain Tumor Genome Anatomy Project

(CGAP/BTGA), Tumor Gene Index

Unpublished (1998)

Other ESTs: 7171d09.x1

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

Tissue Procurement: Robert Jenkins, M.D., Mark Israel, M.D., Jim

Jacobson, Ph.D.

cDNA Library Preparation: David B. Krizman, Ph.D.

DNA Library Arrayed by: Greg Lennon, Ph.D.

Cloning Distribution: Washington University Genome Sequencing Center

found through the I.M.A.G.E. Consortium/LLNL, send email to:

info@image.llnl.gov

Seq primer: -40bp from Gibco.

Location/Qualifiers

1..50

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:3340145"

/clone_lib="NCI_CGAP_Brn20"

/tissue_type="oligodendrogloma"

/dev_stage="adult"

/lab_host="DH10B"

/note="Organ: brain; Vector: pAMP1; mRNA made from

oligodendrogloma tissue, cDNA made by oligo-dT

Directionally cloned. Size-selected on agarose gel,

average insert size 500 bp. Primary library,

non-amplified."

BASE COUNT 33 a 3 c 11 g 3 t

ORIGIN

alignment_scores:

Quality: 45.00 Length: 13

Ratio: 4.091 Gaps: 0

Percent Similarity: 84.615 Percent Identity: 69.231

alignment_block:

US-09-836-410A-1 x BF054800 ..

Align seg 1/1 to: BF054800 from: 1 to: 50

```

344 GluLysGluLysProGlnArgAsnProLysLysLysLys 356
|||||:|||||
2 GAAAAAGAAAGAGGAGGAGGACAAAAA 40

```

seq_name: gb_est1:AA590944

seq_documentation_block:

LOCUS AA590944 50 bp mRNA linear EST 16-SEP-1997
 DEFINITION vm25f02.r1 Knowles Solter mouse blastocyst B1 Mus musculus cDNA
 clone IMAGE:991227 5', mRNA sequence.

ACCESSION AA590944

VERSION AA590944.1 GI:2404257

KEYWORDS

EST.

SOURCE

house mouse.

ORGANISM

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

1 (bases 1 to 50)

Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,

Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,

Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,

Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and

Waterston, R.

The WashU-HMI Mouse EST Project

Unpublished (1996)

Contact: Marra M/Mouse EST Project

WashU-HMI Mouse EST Project

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: mouseest@watson.wustl.edu

This clone is available royalty-free through LLNL; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

MGI:563507.

FEATURES

Location/Qualifiers

1..50

/organism="Mus musculus"

/strain="B6D2 F1/J"

/db_xref="taxon:10090"

/clone="IMAGE:991227"

/clone_lib="Knowles Solter mouse blastocyst B1"

/tissue_type="blastocyst"

/dev_stage="embryo (pre-implantation)"

/lab_host="DH10B"

/note="Organ: embryo; Vector: pSPORT; Site: 1: NOTI;

Site: 2: SalI; Cloned unidirectionally from mRNA prepared

from 800 blastocysts. Primer: SalI(dT):

5'-CGTCGACCGTCGACCGTTTCTTTT-3', cDNAs were

cloned into the NotI/SalI sites of a pSPORT vector (Life

Technologies). Two different size selections: B1 (larger

inserts) and B3."

BASE COUNT 47 a 1 c 1 g 1 t

ORIGIN

alignment_scores:

Quality: 43.00 Length: 16

Ratio: 3.308 Gaps: 0

Percent Similarity: 81.250 Percent Identity: 50.000

alignment_block:

US-09-836-410A-1 x AA590944 ..

Align seg 1/1 to: AA590944 from: 1 to: 50

```

341 LysAsnAlaGluLysGluLysProGlnArgAsnProLysLysLys 356
|||||:|||||
1 AAGAAACAAAAA.....AAAAAAAAAAAAAAAAAAAAA 48

```

```

seq_name: gb_est1:AI813747
seq_documentation_block:
LOCUS      AI813747              37 bp    mRNA          linear      EST 07-MAR-2000
DEFINITION WK79a02.x1 NCI CGAP Panl Homo sapiens cDNA clone IMAGE:2421578 3'
            similar to TR:Q39599 Q39599 EXTENSION; contains element MSRI
            repetitive element ;, mRNA sequence.
ACCESSION  AI813747
VERSION    AI813747.1  GI:5424962
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE  1 (bases 1 to 37)
AUTHORS   NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE     National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
JOURNAL   Unpublished (1997)
COMMENT   Contact: Robert Strausberg, Ph.D.
            Email: cgapbs-r@mail.nih.gov
            Life Technologies catalog #: 11548-013
            DNA Sequencing by: Washington University Genome Sequencing Center
            Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            www-bio.llnl.gov/dbtrp/image/image.html

Trace considered overall poor quality
Insert Length: 718 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 1.
            Location/Qualifiers
FEATURES             source
     1..37
     /organism="Homo sapiens"
     /db_xref="taxon:9606"
     /clone_lib="IMAGE:2421578"
     /clone_lib="NCI CGAP Panl"
     /tissue_type="adenocarcinoma"
     /lab_host="DH10B"
     /note="Organ: pancreas; Vector: pCMV-SPORT6; Site_1: SalI;
     Site_2: NotI; Cloned unidirectionally. Primer: Oligo dT.
     Average insert size 1.72 kb. Life Technologies catalog #:
     11548-013"
BASE COUNT      19 a      16 c      1 g      1 t
ORIGIN
alignment_scores:
    Quality: 42.00      Length: 9
    Ratio: 4.667      Gaps: 0
    Percent Similarity: 100.000      Percent Identity: 77.778
alignment_block:
US-09-836-410A-1 x AI813747 ..
Align seg 1/1 to: AI813747 from: 1 to: 37
348 ProGlnArgAsnProLysLysLys 356
|||||:|||||:|||||:|||||:
6 CCCCCAAAAGCCCCCAAAAAAAAAA 32
seq_name: gb_est2:BM393347
seq_documentation_block:
LOCUS      BM393347              49 bp    mRNA          linear      EST 17-JAN-2002
DEFINITION 50071-2-9-B10.f.1 Chilcoat/Turkewitz cDNA (small fraction)
            Tetrahymena thermophila cDNA, mRNA sequence.
ACCESSION  BM393347
VERSION    BM393347.1  GI:18193400
KEYWORDS   EST.
SOURCE     Tetrahymena thermophila.
            Eukaryota; Alveolata; Ciliophora; Oligohymenophorea;
            Tetrahymena thermophila.
            OrganiSM

```

```

Hymenostomatida; Tetrahymenina; Tetrahymena.
REFERENCE  1 (bases 1 to 49)
AUTHORS   Turkewitz, A.P., Karrer, K.M., Jahn, C., Orlas, E., Kirk, K.E., Frankel
            , J. and Klobutcher, L.
TITLE     EST from Tetrahymena thermophila, strain CU428.1, growing cells
JOURNAL   Unpublished (2002)
COMMENT   Contact: Turkewitz AP
            Molecular Genetics and Cell Biology
            University of Chicago
            920 E. 58th Street, Chicago, IL 60637, USA
            Tel: 773 702 4374
            Fax: 773 702 3172
            Email: apturkew@midway.uchicago.edu
            Seq primer: T3.
            Location/Qualifiers
FEATURES             source
     1..49
     /organism="Tetrahymena thermophila"
     /strain="CU428.1"
     /db_xref="taxon:5911"
     /clone_lib="Chilcoat/Turkewitz cDNA (small fraction)"
     /note="Vector: Bluescript2 SK+; Details on library
     preparation can be found in Chilcoat and Turkewitz (2001)
     Proc. Natl. Acad. Sci USA, 98: 8709-8713."
BASE COUNT      19 a      9 c      7 g      13 t      1 others
ORIGIN
alignment_scores:
    Quality: 41.00      Length: 10
    Ratio: 4.556      Gaps: 0
    Percent Similarity: 90.000      Percent Identity: 50.000
alignment_block:
US-09-836-410A-1 x BM393347/rev ..
Align seg 1/1 to reverse of: BM393347 from: 1 to: 49
236 CysHisGluIleGluArgHisPheIleGlu 245
|||||:|||||:|||||:|||||:
47 TGCCACCGGGTAGAGTGGCATTTTGTCAA 18
seq_name: gb_est2:BM395447
seq_documentation_block:
LOCUS      BM395447              49 bp    mRNA          linear      EST 17-JAN-2002
DEFINITION 50072-2-9-B10.f.1 Chilcoat/Turkewitz cDNA (large fraction)
            Tetrahymena thermophila cDNA, mRNA sequence.
ACCESSION  BM395447
VERSION    BM395447.1  GI:18195500
KEYWORDS   EST.
SOURCE     Tetrahymena thermophila.
            Tetrahymena thermophila
            Eukaryota; Alveolata; Ciliophora; Oligohymenophorea;
            Hymenostomatida; Tetrahymenina; Tetrahymena.
REFERENCE  1 (bases 1 to 49)
AUTHORS   Turkewitz, A.P., Karrer, K.M., Jahn, C., Orlas, E., Kirk, K.E., Frankel
            , J. and Klobutcher, L.
TITLE     EST from Tetrahymena thermophila, strain CU428.1, growing cells
JOURNAL   Unpublished (2002)
COMMENT   Contact: Turkewitz AP
            Molecular Genetics and Cell Biology
            University of Chicago
            920 E. 58th Street, Chicago, IL 60637, USA
            Tel: 773 702 4374
            Fax: 773 702 3172
            Email: apturkew@midway.uchicago.edu
            Seq primer: T3.
            Location/Qualifiers
FEATURES             source
     1..49
     /organism="Tetrahymena thermophila"
     /strain="CU428.1"
     /db_xref="taxon:5911"
     /clone_lib="Chilcoat/Turkewitz cDNA (large fraction)"

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KEYWORDS : house mouse.
SOURCE : Mus musculus
ORGANISM : Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE: 1. (bases 1 to 43)
AUTHORS : Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,
Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly
, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
and Wright, D., Weiss, R.
TITLE : Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
JOURNAL : Unpublished (2000)
COMMENT : Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLCT, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0166 row: E column: 22
Seq primer: CGTTGTAACGACGCGCCAGT
Class: plasmid ends
High quality sequence stop: 43.
Location/Qualifiers
1. .43
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M0166B22"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"
/note="Vector: PWD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(<http://www.jax.org/resources/documents/dnares/>). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (gi14732114|gb|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

```

alignment_scores:
  Quality: 40.00
  Ratio: 4.000
  Percent Similarity: 71.429
  Percent Identity: 50.000
  Gaps: 0
  Length: 14

alignment_block:
  US-09-836-410A-1 x A2860401
  ..

Align seg 1/1 to: A2860401 from: 1 to: 43

341 LysAsnAlaGluLysGluLysProGlnArgAsnProLysLys 354
|||||.....:|||||
2 AAAAACCAACCAACCAACCAACCAACCAACCAACCA 43

seq name: qb_est1:AL587842

```

```

seq_documentation_block:
LOCUS      AL587842      44 bp      mRNA      linear      EST 02-MAR-2001
DEFINITION      BP Chicken Brain Library Gallus gallus cDNA clone
ACCESSION      ROS064C05, mRNA sequence.
VERSION        AL587842
KEYWORDS       EST.
SOURCE         AL587842.1 GI:13192876
              chicken.
ORGANISM       Gallus gallus
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
              Gallinae; Gallus.
REFERENCE      1 (bases 1 to 44)
AUTHORS        Murray, F.
TITLE          BP Chicken Brain Library
JOURNAL        Unpublished (2001)
COMMENT        Contact: Frazer Murray
              Dept. Genomics and Bioinformatics
              Roslin Institute
              Roslin, Midlothian, EH25 9PS, UK
              Tel: +44 (0)131 527 4200
              Fax: +44 (0)131 440 0434
              Email: frazer.murray@bbsrc.ac.uk
              CGCGCGCTTTTCTTTTCTTTTCTTTT 3' Poly A RNA purchased from Clonetech
              (*6854-
              Seq primer: M13F.
              Location/Qualifiers
                source
                  1..44
                  /organism="Gallus gallus"
                  /db_xref="taxon:9031"
                  /clone="ROS064C05"
                  /clone_lib="BP Chicken Brain Library"
                  /tissue_type="Brain"
                  /dev_stage="Unknown"
                  /lab_host="DH10B"
                  /note="Vector: pSPOR1; Site_1: NotI; Site_2: SalI; Cloned
                  unidirectionally. Primer: Oligo dr. 5' adaptor sequence:
                  5' TCGACCTCGAG 3' ; 3' adaptor sequence: 5'
                  CGCGCGCTTTTCTTTTCTTTTCTTTT 3' Poly A RNA purchased from
                  Clonetech (*6854-1)"
BASE COUNT      1 a      0 c      4 g      39 t
ORIGIN
|||||
alignment_scores:
  Quality: 40.00      Length: 12
  Ratio: 3.636      Gaps: 0
  Percent Similarity: 91.667      Percent Identity: 58.333
alignment_block:
  US-09-836-410A-1 x AL587842/rev ..
  Align seg 1/1 to reverse of: AL587842 from: 1 to: 44
  345 LysGLuLysProClnArgAsnProLysLysLysLys 356
  42 AAAAAAAAAAAAAAAAAAACCAAAAAAAAAAAAAA 7
seq_name: gb_est1:AU103783
seq_documentation_block:
LOCUS      AU103783      50 bp      mRNA      linear      EST 30-AUG-2001
DEFINITION      Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
ACCESSION      HEP14796, mRNA sequence.
VERSION        AU103783
KEYWORDS       EST.
SOURCE         AU103783.1 GI:13553304
              human.
ORGANISM       Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 50)

```

```

AUTHORS      Suzuki,Y., Taira,H., Tsunoda,T., Mizushima-Sugano,J., Sese,J., Hata
              ,H., Ota,T., Isogai,T., Tanaka,T., Morishita,S., Okubo,K., Sakaki
              ,Y., Nakamura,Y., Suyama,A. and Sugano,S.
TITLE          Diverse transcriptional initiation revealed by fine, large-scale
              mapping of mRNA start sites
JOURNAL        EMBO Rep. 2 (5), 388-393 (2001)
MEDLINE        21270072
COMMENT        Contact: Yutaka Suzuki
              Department of Virology
              Institute of Medical Science, University of Tokyo
              4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
              Email: ysuzuki@ims.u-tokyo.ac.jp
              Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano
              ,S. Construction and characterization of a full length-enriched and
              a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
FEATURES       Location/Qualifiers
                source
                  1..50
                  /organism="Homo sapiens"
                  /db_xref="taxon:9606"
                  /clone="HEP14796"
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DEFINITION      AU104282 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
ACCESSION      HEP05565, mRNA sequence.
VERSION        AU104282
KEYWORDS       EST.
SOURCE         AU104282.1 GI:13553803
              human.
ORGANISM       Homo sapiens
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              Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 50)
AUTHORS        Suzuki,Y., Taira,H., Tsunoda,T., Mizushima-Sugano,J., Sese,J., Hata
              ,H., Ota,T., Isogai,T., Tanaka,T., Morishita,S., Okubo,K., Sakaki
              ,Y., Nakamura,Y., Suyama,A. and Sugano,S.
TITLE          Diverse transcriptional initiation revealed by fine, large-scale
              mapping of mRNA start sites
JOURNAL        EMBO Rep. 2 (5), 388-393 (2001)
MEDLINE        21270072
COMMENT        Contact: Yutaka Suzuki
              Department of Virology
              Institute of Medical Science, University of Tokyo
              4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
              Email: ysuzuki@ims.u-tokyo.ac.jp
              Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano
              ,S. Construction and characterization of a full length-enriched and
              a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
FEATURES       Location/Qualifiers
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458 GGGACTCGTGCCAGAAAGACTGCCCTTTAACTTTTATCTGGAGAGAAGT 507
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34 heLysGluCysLeuAspArgPheLeuArgMetAsnPheSerLysGlyCys 50
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558 CCACGTGCTCTCAATACCTTGGAGTCTTTATACAGAGATAAAGAGAAGGT 607
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|||||
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2008 AAGCCTTGCTGATGGTATGACCTAGAGACTGTAAGAAGCTGCCGAAGCC 2057
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ID AAH77156 standard; cDNA; 3418 BP.

XX AC AAH77156;

XX DT 21-JAN-2002 (first entry)

XX DE Human tubedown-1 cDNA.

KW Human; tubedown-1; tbdn-1; antisense; cytostatic; osteopathic;
KW bone tumour; osteosarcoma; Ewings sarcoma; metastasis; ss.

XX OS Homo sapiens.

XX FH Key Location/Qualifiers
XX CDS 408..2189

XX FT /*tag= a
XX FT /product= *tubedown-1*

XX PN WO200179505-A2.

XX PD 25-OCT-2001.

XX PE 17-APR-2001; 2001WO-US12435.

XX PR 17-APR-2000; 2000US-197977P.

XX PR 17-APR-2001; 2001US-0836410.

XX PA (CHIL-) CHILDRENS HOSPITAL RES FOUND.

XX PI Gendron RL, Paradis H;

XX DR WPI; 2002-017618/02.

XX DR P-PSDB; AAG77907.

XX PT Nucleic acid molecules antisense to the tubedown-1 gene prevent
XX PT overexpression of tubedown-1 protein and are useful to treat
XX PT osteosarcoma and Ewing's Sarcoma family of tumours

XX PS Claim 1; Page 36-38; 56pp; English.

XX

CC The sequence represents a new human gene, tubedown-1 (tbdn-1). The
CC invention relates to a novel isolated nucleic acid of the tubedown-1
CC gene, and antisense nucleic acids to tbdn-1. The polynucleotides and
CC protein of the invention have cytostatic and osteopathic activity. The
CC polynucleotides of the invention may be used in antisense-therapy/gene
CC therapy. They are useful in the treatment of bone tumours, especially
CC osteosarcoma and Ewings sarcoma family of tumours. The compounds of the
CC invention may also be useful for the prevention of metastases from these
CC types of tumours, either alone or in combination with radiotherapy and/or
CC chemotherapeutic agents.

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Ratio: 5.234 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 100.000

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US-09-836-410A-1 x AAH77156 ..

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51 ProProValPheAsnThrLeuArgSerLeuTyrArgAspLysGluLysVa 67
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858 AATATTAAAGAAAGCTCCAGGTGGATGGTGAAGCCAGGCCCTGGACAC 907
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908 AGCAGACAGATTATTAAATCCAAAGTGTGCAAAATACATGTTAAAAGCCA 957
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958 ACCTGATTAAAGAGGCTGAAGAAATGTCTTCCAGTTTACGAGGAGGA 1007
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XX
AC AAH77158;
XX
DT 21-JAN-2002 (first entry)
XX
DE Human tubedown-1 base pairs 3418-1 antisense cDNA.
XX
KW Human; tubedown-1; tbdn-1; antisense; cytostatic; osteopathic;
KW bone tumour; osteosarcoma; Ewings sarcoma; metastasis; ss.
XX
OS Homo sapiens.
XX
PN WO200179505-A2.
XX
PD 25-OCT-2001.
XX
PF 17-APR-2001; 2001WO-US12435.
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PR 17-APR-2000; 2000US-197977P.
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PR 17-APR-2001; 2001US-0836410.
XX
PA (CHIL-) CHILDRENS HOSPITAL RES FOUND.
XX
PI Gendron RL, Paradis H;
XX
DR WPI; 2002-017618/02.
XX
PT Nucleic acid molecules antisense to the tubedown-1 gene prevent
PT overexpression of tubedown-1 protein and are useful to treat
PT osteosarcoma and Ewing's Sarcoma family of tumours
XX
PS Claim 7; Page 39-41; 56pp; English.
XX
CC The sequence represents tubedown-1 (tbdn-1) bases 3418-1 antisense cDNA.
CC The invention relates to a novel isolated nucleic acid of the tubedown-1
CC gene, and antisense nucleic acids to tbdn-1. The polynucleotides and
CC protein of the invention have cytostatic and osteopathic activity. The
CC polynucleotides of the invention may be used in antisense-therapy/gene
CC therapy. They are useful in the treatment of bone tumours, especially
CC osteosarcoma and Ewings sarcoma family of tumours. The compounds of the
CC invention may also be useful for the prevention of metastases from these
CC types of tumours, either alone or in combination with radiotherapy and/or
CC chemotherapeutic agents.
XX
SQ Sequence 3418 BP; 953 A; 704 C; 604 G; 1157 T; 0 other;

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Quality: 3104.00 Length: 593
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 1511 TATTAGATTCTTCTAGTCAAAAACGAGCAATAGAGCTGGCGACACACT 1462

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Mon Jul 22 09:40:54 2002

us-09-836-410a-1.p2n.rng

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XX AC AAS71925;

XX DT 13-FEB-2002 (first entry)

XX DE DNA encoding novel human diagnostic protein #7729.

XX KW Human; chromosome mapping; gene mapping; gene therapy; forensic;

XX KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

XX OS Homo sapiens.

XX PN WO200175067-A2.

XX PD 11-OCT-2001.

XX PF 30-MAR-2001; 2001WO-US08631.

XX PR 31-MAR-2000; 2000US-0540217.

XX PR 23-AUG-2000; 2000US-0649167.

XX PA (HYSE-) HYSEQ INC.

XX PI Drmanac RT, Liu C, Tang YT;

XX WPI; 2001-639362/73.

XX P-PSDB; ABG07738.

XX PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -

XX PS Claim 1; SEQ ID NO 7729; 103pp; English.

XX CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX SQ Sequence 2477 BP; 859 A; 437 C; 522 G; 659 T; 0 other;

alignment_scores:
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 Ratio: 4.581 Gaps: 16
 Percent Similarity: 82.566 Percent Identity: 80.099
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 1 MetLeuGluArgLeuLysIleThrGluAlaTyrThrLysTyrProAr 17
 535 ATGTTAGAACGGCTAAATAATTTATGAGGAAGCTGGACTAATATATCCAG 584
 17 gGlyLeuValProArgLysLeuProLeuAsnPheLeuSerGlyGluLysP 34
 585 GGGACTGGTGCCCAAGAGCTGCCGTAAACCTTTTATCTGGTGAAGT 634
 34 helysGluCysLeuAspArgPheLeuArgMetAsnPheSerLysGlyCys 50
 635 TTAAGAATGTTGGATAAGTTCTTAAGGATGAATTCAGCAAGGGTTGC 684
 51 ProProValPheAsnThrLeuArgSerLeu.TyrArgAspLysGluLys. 66
 685 CCACCACTCTTCAATACTTTAAGATCATTTACTACCAAGACAAAGAAAGT 734
 67 ValAlaIleValGluGluLeuValValGlyTyrGluThrSerLeuLysSe 83
 735 GTGGCAATCATGAAAGTTAGTAGTAGTTATGAAACCTCTCTAAAGAG 784
 83 rCysArgLeuPheAsnProAsnAspAspGlyLysGluGluProProThrT 100
 785 CTGCCGGCTATTTAACCCCAATGATGAGAAAGGAGGAAACCAACCA 834
 100 hrLeuLeuTyrValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGly 116
 835 CATTACTTTGGGTCCAGTACTTGTGCACACATTTATGACAAATTTGGT 884
 117 GlnProSerIleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrPr 133
 885 CAGCCATCTATTGCTTTGGAGTACATAAATACTGCTATTGAAAGTACACC 934
 133 oThrLeuIleGluLeuPheLeuValLysAlaLysIleTyrLysHisAlaG 150
 935 TACATTAAATAGAACTCTTTCTCGTGAAGCTAAATCTATAAGCATGCTG 984
 150 lyAsnIleLysGluAlaAlaArgTrpMetAspGluAlaGlnAlaLeuAsp 166
 985 GAAATATTAAGAAAGCTGCAAGGTGGATGGATGAGCCCGCCGCTGGAC 1034
 167 ThrAlaAspArgPheIleAsnSerLysCysAlaLysTyrMetLeuLysAl 183
 1035 ACAGCAGACAGATTTATCAACTCCCAATGTCGCAAAATACATGCTAAAGC 1084
 183 aAsnLeuIleLysGluAlaGluGluMetCysSerLysPheThrArgGluG 200
 1085 CAACCTGATTAAAGAGCTGAAGACATGAGCTCAAGCTTTTACAAGGGAG 1134
 200 lyThrSerAlaValGluAsnLeuAsnGluMetGlnCysMetTrpPheGln 216
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 217 ThrGluCysAlaGlnAlaTyrLysAlaMetAsnLysPheGlyGluAlaLe 233
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 233 uLysLysCysHisGluIleGluArgHisPheIleGluIleThrAspAspG 250
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 250 lnPheAspPheHisThrTyrCysMetArgLysIleThrLeuArgSerTyr 266
 1285 AGTTTGACTTTTCATACATCTGTATGAGGAAGATTACCCTTAGATCATAT 1334

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1335 GTGGACTTATTAAACTAGAGATGACTTCCACAGCATCCATTTCCT 1384
283 eLysAla...AlaArgIleAlaIleGluLeuTyrLeuLysLeuHisAspA 299
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299 snProLeuThrAspGluAsn...LysGluHisGluAlaAspThrAla. 313
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314 AsnMetSerAspLysGluLeuLysLysLeuArgAsnLysGlnArgAla 330
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1485 AACATGCTGACAAAGAGCTAAAGAACGCTACGTAAATAACAAAGAACAGC 1534
330 aGlnLysLysAlaGlnIleGluGluLysLysAsnAlaGluLysGlu 347
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347 ysProGlnArgAsnProLysLysLysAspAspAspGluGluIle 363
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1585 AGCAGCAGAGAAATCACAAGAAAGAGAGATGATGATGAGGAGATA 1634
364 GlyClyProLysGluGluLeuIleProGluLysLeuAlaLysValGluTh 380
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1635 GGAGGTCCAAAGAAAGAACTTATCCAGAGAAAGTGGCCAAAGTTGAAC 1684
380 rProLeuGluAlaIleLysPheLeuThrProLeuLysAsnLeuVal 397
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1685 TCCATTGGAAGAGCTATTAAATTTTAAACACCCGTTGAAGAACTTGTGA 1734
397 ysAsnLysIleGluThrHisLeuPheAlaPheGluIleTyrPheArgLys 413
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447 alCysGluSerLysAspLeuProGluThrValArgThrValLeuLysGln 463
1784 1784
464 GluMetAsnArgLeuPheGlyAlaThrAsnProLysAsnPheAsnGluTh 480
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1792 AAATGGTATTACTATAGATCTCTTCTAGTCAGAAAGCGAGCTATAGAGTTG 1841
514 AlaThrThrLeuAspGlySerLeuThrAsnArgAsnLeuGlnThrCysMe 530
1842 GCAACAACACTTGATGAATCTCTACTACAGAAACCTCCAGACATGTAT 1891
530 tGluValLeuGluAlaLeuCysAspGlySerLeuArgAspCysLysGluA 547
1892 GGAGGTATTGGAAGCCTTGATGATGGTAGCCTAGGAGACTCTAAGAGAG 1941
547 IsAlaGluAlaTyrArgAlaSerCysHisLysLeuPheProTyrAla. 562
1942 CTGCTGAAATTTATAGACAAATTCCTATAGAGCTTTACCCCTTATGCT 1991

563 LeuAlaPhe...MetProProGlyTyrGluGluAspMetLysIleThrV 578
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1992 TTGSCCTTTCCATGCCCGCCCGGATATGAAGAGATATGAAGATCCACAG 2041
578 alAsnGlyAspSerSerAlaGluThrGluGluLeuAlaAsnGluIle 593
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2042 TTAATGGAGATAGTTCTCGAAGCTGAAGAACTGGCCCAATGAAT 2090

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seq_documentation_block:

ID AAH77157 standard; cDNA; 1413 BP.

XX AAH77157;

AC AAH77157;

DT 21-JAN-2002 (first entry)

XX Human tubedown-1 base pairs 1413-1 antisense cDNA.

DE Human; tubedown-1; tbdn-1; antisense; cytostatic; osteopathic;

KW bone tumour; osteosarcoma; Ewings sarcoma; metastasis; ss.

XX Homo sapiens.

XX WO200179505-A2.

XX 25-OCT-2001.

XX 17-APR-2001; 2001WO-US12435.

XX 17-APR-2000; 2000US-197977P.

XX 17-APR-2001; 2001US-0836410.

XX (CHIL-) CHILDRENS HOSPITAL RES FOUND.

XX Gendron RL, Paradis H;

XX WPI; 2002-017618/02.

XX Nucleic acid molecules antisense to the tubedown-1 gene prevent overexpression of tubedown-1 protein and are useful to treat osteosarcoma and Ewing's Sarcoma family of tumours

XX Claim 6; Page 38-39; 56pp; English.

XX The sequence represents tubedown-1 (tbdn-1) bases 1413-1 antisense cDNA. The invention relates to a novel isolated nucleic acid of the tubedown-1 gene; and antisense nucleic acids to tbdn-1. The polynucleotides and protein of the invention have cytostatic and osteopathic activity. The polynucleotides of the invention may be used in antisense-therapy/gene therapy. They are useful in the treatment of bone tumours, especially osteosarcoma and Ewings sarcoma family of tumours. The compounds of invention may also be useful for the prevention of metastases from these types of tumours, either alone or in combination with radiotherapy and/or chemotherapeutic agents.

XX Sequence 1413 BP; 359 A; 314 C; 244 G; 496 T; 0 other;

alignment_scores:

Quality: 1768.00 Length: 335
Ratio: 5.278 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-836-410A-1 x AAH77157/rev ..

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1 MetLeuGluArgLeuLysIleTyrGluAlaIleThrLysTyrProAr 17
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56 CAAGAGCTAAAGAACTCGCTATACAAAGAGAGCTCAAAAGAAAG 7
334 laGln 335
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6 GCCAG 2

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seq_documentation_block:
ID AAH16408 standard; cDNA; 1802 BP.

XX AC AAH16408;
XX DT 26-JUN-2001 (first entry)
XX DE Human cDNA sequence SEQ ID NO:15380.

XX KW Human; primer; detection: diagnosis; antisense therapy; gene therapy; ss.

XX OS Homo.sapiens.

XX PN EP1074617-A2.

XX PD 07-FEB-2001.

XX PF 28-JUL-2000; 2000EP-0116126.

XX PR 29-JUL-1999; 99JP-0248036.

XX PR 27-AUG-1999; 99JP-0300253.

XX PR 11-JAN-2000; 2000JP-0118776.

XX PR 02-MAY-2000; 2000JP-0183767.

XX PR 09-JUN-2000; 2000JP-0241899.

XX PA (HELI-) HELIX RES INST.

XX PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

XX DR WPI; 2001-318749/34.

XX PS Claim 8; SEQ ID 15380; 2537pp + CD ROM; English.

XX CC The present invention describes primer sets for synthesizing 5602
CC full-length cDNAs defined in the specification. Where a primer set
CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
CC to the complementary strand of a polynucleotide which comprises one of
CC the 5602 nucleotide sequences defined in the specification, where the
CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC of an oligonucleotide comprising a sequence complementary to the
CC complementary strand of a polynucleotide which comprises a 5'-end
CC sequence and an oligonucleotide comprising a sequence complementary to a
CC polynucleotide which comprises at least 15 nucleotides and the combination of
CC the 5'-end sequence/3'-end sequence is selected from those defined in
CC the specification. The primer sets can be used in antisense therapy and
CC in gene therapy. The primers are useful for synthesizing polynucleotides,
CC particularly full-length cDNAs. The primers allow obtaining of the full-length
CC detection and/or diagnosis of the abnormality of the proteins encoded by
CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to
CC AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC represent oligonucleotides, all of which are used in the exemplification
CC of the present invention.

XX CC Sequence 1802 BP; 644 A; 298 C; 395 G; 465 T; 0 other;

XX SQ

17 gGlyLeuValProArgLysLeuProLeuAsnPheLeuSerGlyGluLysP 34
1956 GGGACTCGTCCCAAGAAAGCTCCCTTAACTTTATCTGGAGAGAGT 907
34 helysGluCysLeuAspArgPheLeuArgMetAsnPheSerLysGlyCys 50
906 TTAAGGAGTGTTGGATAGTGGTCTCCCTAAGGATGAATTCAGCAAGGCGTGT 857
51 ProProValPheAsnThrLeuArgSerLeuTyrArgAspLysGluLysVa 67
856 CCACCTGTCTTCAATACCTTGGAGTCTTTATACAGAGATAAAGAGAGGT 807
67 lAlaIleValGluGluLeuValValGlyTyrGluThrSerLeuLysSerC 84
806 GGCAATCGTAGAAGAACTAGTAGTTGGTTAAGAACTTCTCTAATAAGTT 757
84 ySArgLeuPheAsnProAsnAspAspGlyLysGluGluProProThrThr 100
756 GTGCCCTATTATACCCCAATGATGATGGAAGGAGGAACCTCCAACCA 707
101 LeuLeuTrpValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyG 117
706 TTACTTTGGGTCCAGTACTATTTGGCACACATATATGATAAATTTGTCA 657
117 nProSerIleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrProt 134
656 GCCATCCATTCTCTGGAATACATAAATACTGCAATTTGAAAGTACACCA 607
134 hrLeuIleGluLeuPheLeuValLysAlaLysIleTyrLysHisAlaGly 150
606 CATTGATAGAACTCTCTCTTAAAGAGCTAAATCTATAAGCATGCTGGG 557
151 AsnIleLysGluAlaAlaArgTrpMetAspGluAlaGlnAlaLeuAspTh 167
556 AATATTAAAGAAAGCTGCCAGGTGGATGATGATGATGATGATGATGATG 507
167 rAlaAspArgPheIleAsnSerLysCysAlaLysTyrMetLeuLysAla 184
506 AGCAGACAGATTATTATTATCCAGTGTGCAAAATACATGTTAAAGCCA 457
184 snLeuIleLysGluAlaGluMetCysSerLysPheThrArgGluGly 200
456 ACCTGATTAAAGAGAGCTCAAGAAATGTGTCCAGTTTACGAGGGAAGA 407
201 ThrSerAlaValGluAsnLeuAlaGlnMetGlnCysMetTrpPheGlnTh 217
406 ACTTCAGCGGTAGAAACCTGTAATGCAATGCAATGCAATGCAATGCAAT 357
217 rGluCysAlaGlnAlaTyrLysAlaMetAsnLysPheGlyGluAlaLeu 234
356 AGAGTGTGCTCAGGCATACAAAGCAATGAACAAATTTGGTGAAGCACTTA 307
234 yLysCysHisGluIleGluArgHisPheIleGluIleThrAspAspGln 250
306 AGAAATGTCTGAAATTTGAGAGACATTTATAGAAATCACCAGTACCAG 257
251 PheAspPheHisThrTyrCysMetArgLysIleThrLeuArgSerTyrVa 267
256 TTTGACTTTTCATACATACATGATGATGAGGAAGATCACCCCTTAGATCAT 207
267 lAspLeuLeuLysLeuGluAspValLeuArgGlnHisPropheTyrPheL 284
206 GGACTTTATTAACCTAGAGATGATCTCTGACAGCATCCATTTTACTTCA 157
284 yAlaAlaArgIleAlaIleGluIleTyrLeuLysLeuHisAspAsnPro 300
156 AAGCAGCGGAGATTTGCTTTTGAAGTCTATTGAGCTTTCATGACACCCCT 107
301 LeuThrAspGluAsnLysGluHisGluAlaAspThrAlaAsnMetSerAs 317
106 CTGACAGATGAGAAACAAAGAACACGAGGCTGATACAGCAACATGCTGA 57
317 pLysGluLeuLysLysLeuArgAsnLysGlnArgArgAlaGlnLysLysA 334

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Percent Similarity: 99.702 Percent Identity: 98.512
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167 rAlaAspArgPheIleAsnSerLysCysAlaLysTyrMetLeuLysAla 184
1294 AGCAGACAGATTTATCACTCCCAATGTGCAAAATACATGCTAAAGCCA 1343
184 snLeuIleLysGluAlaGluLysMetCysSerLysPheThrArgGluGly 200
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201 ThrSerAlaValGluAsnLeuAsnGluMetGlnCysMetTrpPheGln 217
1394 ACATACGGCTAGAGAAATTTGATGAATGCAATGCGATGCTGGTTCCAA 1443
217 rGluCysAlaGlnAlaTyrLysAlaMetAsnLysPheGlyGluAlaLeu 234
1444 AGAATGTGCCAGCTTATAAAGCAATGAATAAATTTGTTGAAGCACTTA 1493
234 ysLysCysHisGluIleGluArgHisPheIleGluIleThrAspGln 250
1494 AGAATGTCCAGAGATGAGACATTTTATAGAAATCACTGATGACCA 1543
251 PheAspPheHisThrTyrCysMetArgLysIleThrLeuArgSerTyrVa 267
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1544 TTGACTTTTCATACATCTGTATGAGGAAGATTACCCCTTAGATCATATGT 1593
267 lAspLeuLeuLysLeuGluAspValLeuArgGlnHisProPheTyrPheL 284
1594 GGACTTATTAAGACTAGAGATGTACTTCGACAGCATCCATTTTACTTCA 1643
284 ysAlaAlaArgIleAlaIleGluIleTyrLeuLysLeuHisAspAsnPro 300
1644 AGCAGCAAGAAATTGCTATAGAGATCTATTGGAAGCTTCATGACAACCC 1693
301 LeuThrAspGluAsnLysGluHisGluAlaAspThrAlaAsnMetSerAs 317
1694 CTTACACATGAGATAAAGAACACGAGCTGATACAGCAACATGCTCTGA 1743
317 pLysGluLeuLysLysLeuArgAsnLysGlnArgAlaGlnLysLysA 334
1744 CAAGAGCTAAAGAGCTACCTATAAACAAGAGAGCTCAAAAGAAAG 1793
334 laGlnIle 336
1794 CCCAGATA 1801
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seq_documentation_block:

ID: AAH16424 standard: cDNA; 1985 BP.

XX AC AAH16424;

XX DT 26-JUN-2001 (first entry)

XX DE Human cDNA sequence SEQ ID NO:15407.

XX KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.

XX OS Homo sapiens.

XX PN EPI074617-A2.

XX PD 07-FEB-2001.

XX PF 28-JUL-2000; 2000EP-0116126.

XX PR 29-JUL-1999; 95JP-0248036.

XX PR 27-AUG-1999; 95JP-0300253.

XX PR 11-JAN-2000; 2000JP-0118776.

XX PR 02-MAY-2000; 2000JP-0183767.

XX PR 09-JUN-2000; 2000JP-0241899.

XX PA (HELI-) HELIX RES INST.

XX PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

XX PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

XX DR WPI; 2001-318749/34.

XX PT Primer sets for synthesizing polynucleotides, particularly the 5602

XX PT full-length cDNAs defined in the specification, and for the detection

XX PT full-length cDNAs -

XX PS Claim 8; SEQ ID 15407; 2537pp + CD ROM; English.

XX CC The present invention describes primer sets for synthesizing 5602

XX CC full-length cDNAs defined in the specification. Where a primer set

XX CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary

XX CC to the complementary strand of a polynucleotide which comprises one of

XX CC the 5602 nucleotide sequences defined in the specification, where the

XX CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination

XX CC of an oligonucleotide comprising a sequence complementary to the

XX CC complementary strand of a polynucleotide which comprises a 5'-end

XX CC sequence and an oligonucleotide comprising a sequence complementary to a

XX CC polynucleotide which comprises a 3'-end sequence, where the

CC oligonucleotide comprises at least 15 nucleotides and the combination of
 CC the 5'-end sequence/3'-end sequence is selected from those defined in
 CC the specification. The primer sets can be used in antisense therapy and
 CC in gene therapy. The primers are useful for synthesizing polynucleotides,
 CC particularly full-length cDNAs. The primers are also useful for the
 CC detection and/or diagnosis of the abnormality of the proteins encoded by
 CC the full-length cDNAs. The primers allow obtaining of the full-length
 CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
 CC AAH13632 to AAH18742 represent human cDNA sequences; AAH92446 to
 CC AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632
 CC represent oligonucleotides, all of which are used in the exemplification
 CC of the present invention.

XX SQ Sequence 1985 BP; 652 A; 349 C; 492 G; 492 T; 0 other;

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 Ratio: 5.175 Gaps: 1
 Percent Similarity: 98.658 Percent Identity: 97.315

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 17 gGlyLeuValProArgLysLeuProLeuAsnPheLeuSerGlyGlyLysP 34
 1146 GGGACTGGTGCCAGAGAGCGCTGCGTTAACTTTTATCTGTCGAGAAGT 1195
 34 heLysGluCysLeuAspArgPheLeuArgMetAsnPheSerLysGlyCys 50
 1196 TTAAAGAATGTTGGATAAGTTCTTCAAGGATGAATTTTCAGCAAGGGTTGC 1245
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 1246 CCACCAGCTCTTCAATACCTTTTATGATCATTTACAAAGACAAAGAAAGGT 1295
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 1396 TTACTTTGGTCCAGTACTTCTGGCACAACTATGACAAATTTGGTCA 1445
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 267 lAspLeuLeuLysLeuGluAspValLeuArgGlnHisPropheTyrPheL 284
 1886 GGACTTATTAAACTAGAGATGCTACTTCGACAGCATCCATTTTACTTCA 1935
 284 ysAlaAlaArgIleAlaIleGluIleTyrLeuLysLeuHisasp 298
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seq_documentation_block:
 ID AAH1664 standard; DNA; 710 BP.

XX AC AAH1664;

XX DT 21-SEP-2001 (first entry)

XX DE Human differential transcription-associated cDNA SEQ ID 173.

XX KW Differential transcription; human; rat; tumour cell; cytostatic;

XX KW Ras modulator; Class II tumour suppressor gene; gene therapy; ss.

XX OS Homo.sapiens.

XX PN WO200157058-A2.

XX PD 09-AUG-2001.

XX PF 31-JAN-2001; 2001WO-EP01003.

XX PR 31-JAN-2000; *2000DE-1003102
 XX PA (META-) METAGEN GES GENOMFORSCHUNG MBH.

XX PI Rosenthal A, Hinzmann B, Schaefer R, Zuber J, Tchernitsa O;

XX PI Grips M, Hellriegel M, Schmitz A, Sers C;

XX DR WPI; 2001-483415/52.

XX PT Nucleic acids differentially expressed between tumor and normal cells,
 XX PT useful for diagnosis or therapy of tumors and for screening active
 XX PT agents

XX PS Disclosure; Page 376; 579pp; German.

XX CC This invention describes a nucleic acid (I) with differential expression
 CC between tumour and normal cells and which has cytostatic activity. (I)
 CC work as modulators of Ras activity by inducing expression of tumour
 CC suppressor genes. (I), and polypeptides encoded by them, are useful as
 CC targets for diagnosis or therapy and in screening to determine the
 CC effects of an active compound (potential pharmaceutical) on a cell line,
 CC particularly for diagnosis and treatment of tumors, especially by
 CC modulating expression of (I) (by gene therapy, antisense RNA or ribozyme
 CC methods) or by modulating the amount and/or location of (I)-encoded
 CC polypeptides (by administration of the polypeptide or its activator,
 CC antibody (optionally as a conjugate) or inhibitor). The method allows

CC Identification of many Class II tumour suppressor genes (i.e. genes that
 CC are not primary targets for tumour-initiating mutations).
 CC AAH81492-AAH82376 represent the human and rat derived nucleic acid
 CC fragments described in the method of the invention.
 SQ Sequence 710 BP; 170 A; 146 C; 121 G; 267 T; 6 other;

alignment_scores:
 Quality: 1176.00 Length: 237
 Ratio: 5.091 Gaps: 0
 Percent Similarity: 97.468 Percent Identity: 96.203

alignment_block:
 US-09-836-410A-1 x AAH81664/rev ..

Align seg 1/1 to reverse of: AAH81664 from: 1 to: 710

```

103 TrpValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyGlnProse 119
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
710 TGGGTCAAGTACTTGGCACACCATTTATGACAAAATTGGTCAGCCATC 661
119 rIleAlaLeuGluTyrIleAsnThrAlaIleGluSerThrProThrLeuI 136
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
660 TANTGCTNTGGAGTACATAAATCTGCTATTGANAGTACACT ACATTA 612
136 leGluLeuPheLeuValLysAlaLysIleTyrLysHisAlaGlyAsnIle 152
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
611 TAGACNTCTTCTCGTGANAGCTAAATCTATAAGCATGCTGGAATATT 562
153 LysGluAlaAlaArgTrpMetAspGluAlaGlnAlaLeuAspThrAlaAs 169
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
561 AAAGAAGCTGCAAGTGGATGGATGAGCNCAGGCCCTTGGACACACAGA 512
169 pArgPheIleAsnSerLysCysAlaLysTyrMetLeuLysAlaAsnLeuI 186
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
511 CAGATTATCAACTCCAAATGTGCAAAATACATGCTAAAAGGCCAAGCTGA 462
186 leLysGluAlaGluMetCysSerLysPheThrArgGluGlyThrSer 202
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
461 TTAAGAAGCTGAAGAAATGTCTCAAACTTACAAAGGAAGGACATCA 412
203 AlaValGluAsnLeuAsnGluMetGlnCysMetTrpPheGlnThrGluCy 219
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
411 GCGGTAGAGAAATTTGAATGAAATGCGATGCTGTTCCAAACAGCAATG 362
219 sAlaGlnAlaTyrLysAlaMetAsnLysPheGlyGluAlaLeuLysLysC 236
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
361 TGCCCAAGCTTATTAAGCAATGAATAATTTGGTGAAGCACCTTAAGAAAT 312
236 yHisGluIleGluArgHisPheIleGluIleThrAspAspGlnPheAsp 252
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
311 GTCATGAGATTGAGAGACATTTTATAGAAATCACTGATGACCATGTTGAC 262
253 PheHisThrTyrCysMetArgLysIleThrLeuArgSerTyrValAspLe 269
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
261 TTTTACATACATCTGTATGAGGAAGATTACCTCTAGATCATATGTGACTT 212
269 uLeuLysLeuGluAspValLeuArgGlnHisProPheTyrPheLysAlaA 286
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
211 ATTAAGCTAGAGATGCTCTCGACACATCCATTTTACTTCAAGGGCAG 162
286 laArgGlaAlaIleGluIleTyrLeuLysLeuHisAspAsnProLeuThr 302
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
161 CAAGAATTGCTATAGATGATCTATTTGAAGCTTCATGACAAACCCCTTACA 112
303 AspGluAsnLysGluHisGluAlaAspThrAlaAsnMetSerAspLysG 319
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
111 GATGAGATAAAGAACACGAAGCTGATACAGCAACATGTCTGACAAAGA 62
319 uLeuLysLysLeuArgAsnLysGlnArgAlaGlnLysLysAlaGlnI 336
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
61 GCTAAGAAGCTACGTATATAACAAAGAGAGCTCAAAAGAAAGCCCAAGA 12

```

336 leGluGluGlu 339
 |||||:|||||
 11 TAGAAGAAGAG 1

seq_name: /SIDSI/cgdata/hold-geneseq/geneseqn-emb1/NA1999.DAT:AAZ15705

seq_documentation_block:

XX ID AAZ15705 standard; cDNA; 781 BP.

XX AC AAZ15705;

XX DT 12-OCT-1999 (first entry)

XX DE Human gene expression product cDNA sequence SEQ ID NO:3174.

XX KW Human; gene: gene expression product; diagnosis; therapy; probe;

XX KW detection; mapping; tissue typing; profiling; forensic; cancer;

XX KW genetic analysis; colorectal cancer; breast cancer; lung cancer; ss.

XX OS Homo sapiens.

XX PN WO9938972-A2.

XX PD 05-AUG-1999.

XX 28-JAN-1999; 99WO-US01619.

XX 03-APR-1998; 98US-0080666.

XX 28-JAN-1998; 98US-0072910.

XX 24-FEB-1998; 98US-0075954.

XX 31-MAR-1998; 98US-0080114.

XX 03-APR-1998; 98US-0080515.

XX (CHIR) CHIRON CORP.

XX (HYSE-) HYSEQ INC.

XX Crkvenjakov R, Dickson M, Drmanac R, Drmanac S;

XX Escobedo J, Garcia PD, Garcia V, Glese K, Innis MA;

XX Jones WL, Kassam A, Kennedy GC, Kita D, Labat I;

XX Lamson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;

XX Stache-Crain B, Sudduth-Klinger J, Williams LT;

XX WPI; 1999-494092/41.

XX Novel human genes and their expression products which are
 PT differentially expressed in different cell types

XX Claim 1; Page 1524-1525; 2479pp; English.

XX The present invention describes a library of human polynucleotides
 CC comprising the sequences given in AAZ1532 to AAZ1779. Also described is
 CC a method of detecting differentially expressed genes correlated with the
 CC cancerous state of a mammalian cell, comprising detecting at least one
 CC differentially expressed gene product in a test sample from a cell
 CC suspected of being cancerous, where the gene product is encoded by one
 CC of the 5248 polynucleotide sequences given in AAZ1532 to AAZ1779. The
 CC polynucleotides can be used as a source of primers and probes, which can
 CC be used for a variety of purpose, e.g. detection of expression levels,
 CC mapping, tissue typing or profiling, forensics, genetic analysis and
 CC detection of polymorphisms. Polypeptides encoded by the polynucleotides
 CC can be used for raising antibodies for experimental, diagnostic and
 CC therapeutic purposes. The polynucleotides may also be used to construct
 CC arrays for diagnostics (which may be used to determine function of an
 CC encoded protein); and to detect differences in expression levels between
 CC two cells (e.g. to identify abnormal or diseased tissue in a human, to
 CC identify a genetic predisposition or susceptibility to a disease such as
 CC cancer). The polynucleotides of the invention are especially used in the
 CC diagnosis, prognosis and management of colorectal cancer, breast cancer,
 CC and lung cancer. The polynucleotides can also be used to screen for
 CC peptide analogues and antagonists.

XX Sequence 781 BP; 263 A; 140 C; 156 G; 205 T; 17 other;

seq_name: /SIDS1/gcgdata/hold-geneseq/genesequ-emb1/NA1999.DAT: AAX99053
seq_documentation_block:
ID AAX99053 standard; cDNA; 781 BP.
XX

CC mammary dysplasia, hyperplasias, e.g. endometrial, adrenal, breast,
CC prostate or thyroid hyperplasias or pseudoeplitheliomatous hyperplasia of
CC the skin.
XX
SQ Sequence 781 BP; 263 A; 140 C; 156 G; 205 T; 17 other;

alignment_scores:
Quality: 1104.00 Length: 232
Ratio: 4.973 Gaps: 4
Percent Similarity: 95.690 Percent Identity: 95.259

alignment_block:

US-09-836-410A-1 x AAX99053 ..

Align seg 1/1 to: AAX99053 from: 1 to: 781

62 ArgAspLysGlu.LysValAlaLeValGluLeuValValcylTyrG 78
87 AGAGACAAAGAAAAGGTGCAATCATAGAGAGATTAGTAGGTTATG 136
78 luThrSerLeuLysSerCysArgLeuPheAsnProAsnAspGlyLys 94
137 AACCTCTCTAAAAGCTGCGGTATTAAACCCCAATGATGAAAG 186
95 GluGluProProThrThrLeuLeuTrpValGlnTyrTyrLeuAlaGlnHi 111
187 GAGAACCAACCAACCATACATTCTGGTCCNNTCTACTTGGCACAACA 236
111 sTyrAspLysIleGlyGlnProSerIleAlaLeuGluTyrIleAsnThra 128
237 TTATGACAAATTTGGTCAGCCATCTATTGCTTTGGAGTACATAAATCTG 286
128 laileGluSerThrProThrLeuIleGluLeuPheLeuValLysAlaLys 144
287 CTATTGAAGTACACCTACATTAATAGACTCTTCTCGTGAAGTAA 336
145 IleTyrLysHisIleGlyAsnIleLysGluAlaAlaArgTrpMetAspGI 161
337 ATCTATAAGCATGCTGGAATATTAAGAAGCTGCAAGTGGATGATCA 386
161 uAlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSerLysCysAlaL 178
387 GGCACAGGCTTGGACACAGACAGATTTATCAACTCCAAATGTGCAA 436
178 ysTyrMetLysAlaAsnLeuIleLysGluAlaGluLeuMetCysSer 194
437 AATACATGCTAAAGGCAACCTGATTAAAGAGCTGAAGAAATGTGCTCA 486
195 LysPheThrArgGluClyThrSerAlaValGluAsnLeuAsnGluMetGI 211
487 AAGTTTACAGGGAAGGAACATCAGCGGTAGAGAATTTGAATGAATGCA 536
211 nCysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLysAlaMetAsnL 228
537 GTGCATGTGTTCCAAACAGAAATGTCCCGAGCTTATAAGCAATGAATA 586
228 ysPheClyGluAlaLeuLysLysCysHisGluIleGluArgHisPheIle 244
587 AATTTGGTGAAGCACTTAAGAAATCTCATGAGATTGAGAGACATTTATA 636
245 GluIleThrAspAspGlnPheAspPheHisThrTyrCysMet.ArgLys 260
637 GGAATCACTGATGACCACTTTGACTTTTCATACATCTACTGGATGAAGGA 686
261 IleThrLeuArgSerTyrValAspLeuLeuLysLeu.GluAspValLeuA 277
687 ATTACCCCTTAGATCATATGTGGACTTATTNAACTATGAGATGACTTT 736
277 rgGlnHisProPheTyrPheLysAlaAlaArgIleAla 289
737 NACAGCATNCATTTTACTTCAAGGACGACGAAGATTTGCT 774

seq_name: /SIDS1/gcgcdata/hold-geneseq/geneseqn-embl/NA1999.DAT:AAZ15983
seq_documentation_block:
ID AAZ15983 standard; CDNA; 764 BP.
XX
AC AAZ15983;
XX
DT 12-OCT-1999 (first entry)
XX
DE Human gene expression product cDNA sequence SEQ ID NO:3452.
XX
KW Human; gene; gene expression product; diagnosis; therapy; probe;
KW detection; mapping; tissue typing; profiling; forensic; cancer;
KW genetic analysis; colorectal cancer; breast cancer; lung cancer; ss.
XX
OS Homo sapiens.
XX
PN WO9938972-A2.
XX
PD 05-AUG-1999.
XX
PF 28-JAN-1999; 99WO-US01619.
XX
PR 03-APR-1998; 98US-0080666.
PR 28-JAN-1998; 98US-0072910.
PR 24-FEB-1998; 98US-0075954.
PR 31-MAR-1998; 98US-0080114.
PR 03-APR-1998; 98US-0080515.
XX
PA (CHIR) CHIRON CORP.
PA (HYSE-) HYSEQ INC.
XX
PI Crkvenjakov R, Dickson M, Drmanac R, Drmanac S;
PI Escobedo J, Garcia PD, Garcia V, Giese K, Innis MA;
PI Jones WL, Kassam A, Kennedy GC, Kita D, Labat J;
PI Lamson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;
PI Stache-Crain B, Sudduth-Klinger J, Williams LT;
XX
DR WPI; 1999-494092/41.
XX
PT Novel human genes and their expression products which are
PT differentially expressed in different cell types
XX
PS Claim 1; Page 1650-1651; 2479pp; English.
XX

The present invention describes a library of human polynucleotides comprising the sequences given in AAZ12532 to AAZ1779. Also described is a method of detecting differentially expressed genes correlated with the cancerous state of a mammalian cell, comprising detecting at least one differentially expressed gene product in a test sample from a cell suspected of being cancerous, where the gene product is encoded by one of the 5248 polynucleotide sequences given in AAZ12532 to AAZ1779. The polynucleotides can be used as a source of primers and probes, which can be used for a variety of purpose, e.g. detection of expression levels, mapping, tissue typing or profiling, forensics, genetic analysis and detection of polymorphisms. Polypeptides encoded by the polynucleotides can be used for raising antibodies for experimental, diagnostic and therapeutic purposes. The polynucleotides may also be used to construct arrays for diagnostics (which may be used to determine function of an encoded protein); and to detect differences in expression levels between two cells (e.g. to identify abnormal or diseased tissue in a human, to identify a genetic predisposition or susceptibility to a disease such as cancer). The polynucleotides of the invention are especially used in the diagnosis, prognosis and management of colorectal cancer, breast cancer, and lung cancer. The polynucleotides can also be used to screen for peptide analogues and antagonists.

Sequence 764 BP; 253 A; 138 C; 148 G; 204 T; 21 other;

alignment_scores:
Quality: 973.00 Length: 200
Ratio: 5.015 Gaps: 3

XX Cancer; human; colon; breast; lung; transmembrane receptor; ATPase;
 integral membrane protein; aspartyl protease; GARA family; wnt family;
 KW transcription factor; G-protein alpha subunit; protein phosphatase;
 KW phorbol ester binding protein; diacylglycerol binding protein; trypsin;
 KW protein kinase; tyrosine phosphatase; developmental signalling protein;
 KW WW/rsp5/WWP domain; therapy; forensic; genetic mapping; diagnostic;
 KW detection; treatment; cervical; melanoma; colorectal adenocarcinoma;

US-09-836-410A-1 x AAX98777

Align seg 1/1 to: AAX98777 from: 1 to: 764

```
62 ArgAspLysGluLysValAlaIleValGluLeuValValGlyTyrGI 78
|||||
65 AGAGACAAGAAAGGTGGCAATCATAGAGAGTTTNTAGTAGTATGA 114
|||||
78 uThrSerLeuLysSerCysArgLeuPheAsnProAsnAspAspGlyLysG 95
|||||
115 AACCTCTCTAAAGAGTCCCGGTTATTAAACCCCAATCATGATGGAAGG 164
|||||
95 luGluProThrThrLeuLeuTyrValGlnTyrTyrLeuAlaGlnHis 111
|||||
165 AGGAACCAACCAACACATTAATTTGGGTCCAGTACTTGGGCACACAT 214
|||||
112 TyrAspLysIleGlyGlnProSerIleAlaLeuGluTyrIleAsnThrAl 128
|||||
215 TATGACAAATTTGGTCAGCCATCTATTGCTTTGGAGTACATAAATACTGC 264
|||||
128 aileGluSerThrProThrLeuIleGluLeuPheLeuValLysAlaLysI 145
|||||
265 TATTGAAGTACACTACATTAATAGAACTCTTTCTCGTGAAGCTAAAA 314
|||||
145 leTyrLysHisAlaGlyAsnIleLysGluAlaAlaArgTyrMetAspGlu 161
|||||
315 TCTATAAGCATGCTGGAAATATTAAAGAACGTCGAAGTGGATGGATGAG 364
|||||
162 AlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSerLysCysAlaLy 178
|||||
365 GCCCAGGCCCTTGGACACAGACAGAGATTTATCAACTCCAAATGTGCAA 414
|||||
178 sTyrMetLeuLysAlaAsnLeuIleLysGluAlaGluMetCysSerL 195
|||||
415 ATACATGCTAAAGCCAACTGATTAAAGAGCTGAAGAATGTGCTCAA 464
|||||
195 ysPheThrArgGluGlyThrSerAlaValGluAsnLeuAsnGluMetGln 211
|||||
465 AGTTTACAAGGGAAGGAACATCAGCGGTAGAGAAATTTGAATGAATGCAG 514
|||||
212 CysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLysAlaMetAsn.L 228
|||||
515 TGATGTGGTTCCAAACAGAAATGTGCCAGGCTTATAAAGCAATGAATTA 564
|||||
228 ysPheGlyGluAlaLeuLysLysCysHis.GluIleGluArgHisPheI 244
|||||
565 AATTGTGTGAAGCACTTAAGAATGTGATTCAGATTGAGAGACTTTTATA 614
|||||
244 eGluIleThrAspAsp.GlnPheAspPheHisThrTyrCysMet 258
|||||
615 GGAATCACTGATGACCCAGTTTGACTTTTCATACATACATCTATG 658
|||||
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seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-emb1/NA2001A.DAT.AAS37350

seq_documentation_block:

ID AAS37350 standard; cDNA; 402 BP.

XX

AC AAS37350;

XX

DT 17-DEC-2001 (first entry)

XX

DE Novel human diagnostic and therapeutic gene #408.

XX Human; cancer; breast; lung; colon; prostate; cytostatic; diagnostic; ss.

XX Homo sapiens.

OS WO2001:66753-A2.

XX

PD 13-SEP-2001.

XX

PF 09-MAR-2001; 2001WO-US07787.

XX

PR 09-MAR-2000; 2000US-0188609.

XX (CHIR) CHIRON CORP.

PA (HYSE-) HYSEQ INC.

XX Williams LT, Escobedo J, Innis MA, Garcia PD, Sudduth-Klinger J;

PI Reinhard C, Randazzo F, Kennedy GC, Pot D, Kassam A, Lamson G;

PI Drmanac R, Crkvenjakov R, Dickson M, Drmanac S, Lebat I;

PI Leshkowitz D, Kita D, Garcia V, Jones WL, Stache-Crain B;

XX WPI; 2001-530177/58.

DR

XX New polynucleotides and polypeptides, useful for diagnosis and

PT treatment of breast, lung and colon cancer -

XX Claim 1; Page 698-699; 1193pp; English.

XX The invention relates to new polynucleotides and polypeptides, useful for diagnosis and treatment of breast, lung and colon cancer. The sequences can be used in detecting differentially expressed genes correlated with a cancerous state of a mammalian cell, comprising detecting at least one differentially expressed gene product in a test sample derived from a cell suspected of being cancerous. They can also be used to inhibit tumour growth by modulating expression of a gene product. AAS36943-AAS39338 represent novel human diagnostic and therapeutic coding sequences of the invention.

XX Sequence 402 BP; 142 A; 76 C; 85 G; 99 T; 0 Other;

alignment_scores:

Quality: 580.00 Length: 130

Ratio: 5.231 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-836-410A-1 x AAS37350 ..

Align seg 1/1 to: AAS37350 from: 1 to: 402

71 GluGluLeuValValGlyTyrGluThrSerLeuLysSerCysArgLeuPh 87

|||||

12 GAAGAGTAGTAGTAGTATGAAACCTCTCTAAAGAGTCCCGGTTATT 61

|||||

87 eAsnProAsnAspAspGlyLysGluGluProProThrThrLeuLeuTpv 104

|||||

62 TAACCCCAATGATGATGGAAGGAGGAGAACCAACCAACACATTACTTTGGG 111

|||||

104 aGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyGlnProSerIle 120

|||||

112 TCCAGTACTACTTGGCACAACATTATGACAAATTTGGTCAGCCATCTATT 161

|||||

121 AlaLeuGluTyrIleAsnThrAlaIleGluSerThrProThrLeuIleGl 137

|||||

162 GCTTTGGAGTACATAAATACTGCTATTGAAAGTACACCTACATTAATA 211

|||||

137 uLeuPheLeuValLysAlaLysIleTyrLysHisAlaGlyAsnIleLysG 154

|||||

212 ACTCTTCTCGTGAAGCTAAATCTATAAGCATGCTGGAATAATTAAG 261

|||||

154 luAlaAlaArgTyrMetAspGluAlaGlnAlaLeuAspThrAlaAspArg 170

|||||

262 AAGCTGCAAGTGGATGGATGAGCCAGCCCTTGGACACAGACAGACAGA 311

|||||

171 PheIleAsnSerLysCysAlaLysTyrMetLeuLysAlaAsnLeuIleLy 187

|||||

312 TTTATCAACTCCAAATGTGCAAAATACATGCTAAAGCCACCTGATTAA 361

|||||

187 sGluAlaGluMetCysSerLysPheThrArgGluGly 200

|||||

362 AGAAGCTGAAGAAATGTCTCAAAAGTTTACAAGGGAAGGA 401

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-emb1/NA2001A.DAT.AAH12222

Mon Jul 22 09:40:54 2002

us-09-836-410a-1.p2n.rng

Align seq 1/1 to reverse of: AAH12222 from: 1 to: 488

142 LysAlaLysIleTyrLysHisAlaGlyAsnIleLysGluAlaAlaArgTr 158
466 GAAGCTAAATCTATAGCA.CCTGNAANTTTNAAAGAGCCCTNCAAGGG 418
158 pMetAspGluAlaGlnAlaLeuAspThrAlaAspArgPheIleAsnSerL 175
417 GATGGATTAGGCCAGCCGCTTGGACCCAGCAGCCAGTTTATCAACTCCA 368
175 yCysAlaLysTyrMetLeuLysAlaAsnLeuLysGluAlaGluGlu 191
367 AATGTCAAATACATGCTAAAGCCCACTGATTAAAGAAAGCTGAAGA 318
192 MetCysSerLysPheThrArgGluGlyThrSerAlaValGluAsnLeuAs 208
317 ATGTGCTCAAGTTTNCAGGGAAGAACATCAGCGGTANAGAAATTGAA 268
208 nGluMetGlnCysMetTrpPheGlnThrGluCysAlaGlnAlaTyrLysA 225
267 TGAATGTCAGCATGNGGTTCCAAACAGAAATNGCCAGGCTTATAAG 218
225 laMetAsnLysPheGlyGluAlaLeuLysLysCysHisGluIleGluArg 241
217 CAATGAATAAATTTGGTGAAGCNCCTTAAGAAATGTCATGAGATTGAGA 168
242 HisPheIleGluIleThrAspAspGlnPheAspPheHisThrTyrCysMe 258
167 AA.....ATCNCGTGATGACCACTTTGACTTTTCATACATACTGTAT 128
258 tArgLysIleThrIleuArgSerTyrValAspLeuLeuLysLysGluAspV 275
127 GAGGAGATTACCTTATAGATCATATNGGACTTATTAAAACTAGAAGATG 78
275 alLeuArgGlnHisProPheTyrPheLysAlaAlaArgIleAlaIleGlu 291
77 NACTTCGACAGCATCCNTTTTACTTCAAGGCAGCAAGAAATTGCTATANAG 28

292 ileTyrLeuLysLysHisAsp 298
27 ATCTATTGAAGCTTCATGAC 7

seq_name: /SIDS1/gcgdata/hold-geneseq/geneseq-emb1/NA1989.DAT.AAN90541

seq_documentation_block:
ID AAN90541 standard; DNA; 2703 BP.

XX AC AAN90541;
XX DT 28-NOV-1989 (first entry)
XX DE DNA encoding N-alpha-acetyl transferase.
XX KW N-alpha-acetyl transferase; herbicide resistance;
XX KW protein N-acetylation.
XX FH Key Location/Qualifiers
FT misc_feature 272..335 /*tag= a
FT misc_feature 338..392 /*tag= b
FT misc_feature 479..515 /*tag= c
FT misc_feature 542..566 /*tag= d
FT misc_feature 971..989 /*tag= e
FT misc_feature 1007..1049 /*tag= f
FT misc_feature 1061..1085 /*tag= g
FT misc_feature 1088..1130

seq_documentation_block:
ID AAH12222 standard; cDNA; 488 BP.
AC AAH12222;
XX DT 26-JUN-2001 (first entry)
XX DE Human cDNA clone (3'-primer) SEQ ID NO:9057.
XX KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX OS Homo sapiens.
XX PN EP1074617-A2.
XX PD 07-FEB-2001.
XX PF 28-JUL-2000; 2000EP-0116126.
XX PR 29-JUL-1999; 99JP-0248036.
XX PR 27-AUG-1999; 99JP-0300253.
XX PR 11-JAN-2000; 2000JP-0118776.
XX PR 02-MAY-2000; 2000JP-0183767.
XX PR 09-JUN-2000; 2000JP-0241899.
XX PA (HELI-) HELIX RES INST.
XX PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
XX PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX DR WPI: 2001-318749/34.
XX PT Primer sets for synthesizing polynucleotides, particularly the 5602
XX PT full-length cDNAs defined in the specification, and for the detection
XX PT and/or diagnosis of the abnormality of the proteins encoded by the
XX PT full-length cDNAs -
XX PS Claim 3; SEQ ID 9057; 2537pp + CD ROM; English.

XX The present invention describes primer sets for synthesizing 5602
XX full-length cDNAs defined in the specification. Where a primer set
XX comprises: (a) an oligo-dr primer and an oligonucleotide complementary
XX to the complementary strand of a polynucleotide which comprises one of
XX the 5602 nucleotide sequences defined in the specification, where the
XX oligonucleotide comprises at least 15 nucleotides; or (b) a combination
XX of an oligonucleotide comprising a sequence complementary to the
XX complementary strand of a polynucleotide which comprises a 5'-end
XX sequence and an oligonucleotide comprising a sequence complementary to a
XX polynucleotide which comprises a 3'-end sequence, where the
XX oligonucleotide comprises at least 15 nucleotides and the combination of
XX the 5'-end sequence/3'-end sequence is selected from those defined in
XX the specification. The primer sets can be used in antisense therapy and
XX in gene therapy. The primers are useful for synthesizing polynucleotides,
XX particularly full-length cDNAs. The primers are also useful for the
XX detection and/or diagnosis of the abnormality of the proteins encoded by
XX the full-length cDNAs. The primers allow obtaining of the full-length
XX cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
XX AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
XX AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
XX represent oligonucleotides, all of which are used in the exemplification
XX of the present invention.

SQ Sequence 488 BP; 121 A; 98 C; 84 G; 167 T; 18 other;

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Ratio: 4.388 Gaps: 1
Percent Similarity: 87.898 Percent Identity: 82.166
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US-09-836-410A-1 x AAH12222/rev ..

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XX      PD
XX      10-AUG-1989.
XX      PF      07-FEB-1989;      89WO-US00471.
XX      PR      08-FEB-1988;      88US-0153361.
XX      PR      14-DEC-1988;      88US-0284344.
XX      PA      (GEOH ) THE GENERAL HOSPITAL CORPORATION.
XX      PI      Smith JA, Lee FJS;
XX      DR      WPI; 1989-249008/34.
XX      DR      P-PSDB; AAP91070.
XX      PT      New pure N-alpha-acetyl transferase and DNA encoding it - catalysing
XX      PT      acetylation of proteins and peptides, eg to stabilise pharmaceuticals
XX      PT      or induce herbicide resistance in plants.
XX      PS      Claim 8: Page 50; fig 12b-e; 72pp; English.
XX      CC      DNA encodes N-alpha-actyl transferase, used for catalysing N-acetylation
XX      CC      of peptides/proteins, eg to stabilise pharmaceuticals or to induce
XX      CC      herbicide resistance in plants. Features a - n are fragments resulting
XX      CC      from exonuclease III deletion. See also AAP91070.
XX      SQ      Sequence 2703 BP; 943 A; 489 C; 530 G; 741 T; 0 other;

alignment_scores:
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|||||
31 Y...GluLysPheLysGluCysLeuAspArgPheLeuArgMetAsnPhes 47
|||||
963 CAAGAAGAGCTCAGCAAAAATTCAGAGATATGTTTGCCTCAATTGG 1012
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47 erLysGlyCysProValPheAsnThrLeuArgSerLeuTyr...Arg 62
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1013 AGCGGGGTTCAGCAACCTTTTCCAAAGTGAACCCCTTTACCAAGA 1062
|||||
63 AspLysGluLysValAla...IleValGluGluLeuValValGlyTyrG1 78
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1063 AGCAAGTCCAGGTTTCACCACTATTTGGAGAAAATTTGCTTGTATTATT 1112
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1113 GTCCGGATTA.....GATCCTACGCGGAT..... 1137

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95 luGluProProThrThrLeuLeuTirpValGlnTyrTyrLeuAlaGlnHis 111
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1138 .....CCAAATCCTTTTATTTGGACCAANTATTACTTGCTCAACAT 1179
|||||
112 TyrAspLysIleGlyGlnProSerIleAlaLeuGluTyrIleAsnThrAl 128
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1180 TTCCTTTTCTTAAGGATTTTCGAAAGCCCAAGAAATATATTGATGCTGC 1229
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1230 CCTTGACCACACCCCACTTTAGTTGAGTTTACATCTCCCAAGGCAGTA 1279
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Mon Jul 22 09:40:54 2002

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2094 ATTTATAATAACTACTCCATCGCAAGTCAGAGAAGACGAAAGGGATTATA 2143
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436 uHisGluCysMetIleArgLeuPheHisSerValCysGluSerLysAspL 453
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2244 TGGTAGTAGGCCCATTTGTTTGTACATGCCACAAGAAACGACACCCCT 2293
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Date: Jul 20, 2002 3:43 AM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 CompuGen Ltd.

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Query length: 593

Database: GenEmbl.*

Database sequences: 1797656

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gb_pat:AX285242	+ 3104.00	4358.66	2.0e-234	3418	AX285242 Sequence 1 from Patent
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seq_documentation_block:

LOCUS AX285247 1779 bp DNA linear PAT 20-NOV-2001

DEFINITION Sequence 6 from Patent WO0179506.

ACCESSION AX285247

VERSION AX285247.1 GI:17045931

KEYWORDS human.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (sites)

AUTHORS Gendron,R.L. and Paradis,H.

TITLE Treatment of ocular neovascularization and related diseases

JOURNAL Patent: WO 0179506-A 6 25-Oct-2001;

CHILDREN'S Hospital Research Foundation (US)

FEATURES

Location/Qualifiers

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ORIGIN

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67 lalaTleValGluGluLeuValValGlyTyrGluThrSerLeuLysSerC 84

201 GGCAATCTAGAACACTAGTAGTTGGTTATGAAACTTCTCTAAAAGTT 250

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251 CTCGCCATTTTAAACCCCAATGATGAAAGAGGAGAACCTCCCAACACA 300

101 LeuLeuTrpValGlnTyrTyrLeuAlaGlnHisTyrAspLysIleGlyL 117

301 TTACTTTGGGTCAGTACTATTGGCACAGCATTTATGATAAATTTGGTCA 350

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LOCUS AX285242
DEFINITION Sequence 1 from Patent WO01/79506.

ACCESSION AX285242

VERSION AX285242.1 GI:17045930

KEYWORDS

SOURCE human.

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (sites)

Gendron, R.L. and Paradis, H.

Treatment of ocular neovascularization and related diseases

Patient: WO 0179506-A 1 25-OCT-2001.

Children's Hospital Research Foundation (US)

FEATURES

Location/Qualifiers

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BASE COUNT 1157 a 604 c 704 g 953 t

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Ratio: 5.234 Gaps: 0
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67 lAlaIleValGluLeuValValGlyTyrGluThrSerLeuLysSerC 84
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658 GTGCGCTATTATTAACCCCAATGATGTGGAAGAGGAGGAACCTCCAACACA 707
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DEFINITION Sequence 4 from Patent WO01/79505.

ACCESSION AX285296

VERSION AX285296.1 GI:17045977

KEYWORDS human.

SOURCE

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

AUTHORS

TITLE

JOURNAL

INFORMATION

Patent: WO 0179505-A 4 25-OCT-2001;

CHILDREN'S HOSPITAL MEDICAL CENTER (US)

FEATURES

Location/Qualifiers

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Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
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Gendron, R.L., Adams, L.C. and Paradis, H.
Tubedown-1, A novel acetyltransferase associated with blood vessel
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2 (bases 1 to 3421)
Gendron, R.L., Adams, L.C. and Paradis, H.
Direct Submission
Submitted (20-FEB-2000) Pediatrics, Childrens Hospital Medical
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Center, 3333 Burnet Avenue, Cincinnati, OH 45229-3039, USA
Amino acid sequence updated by submitter
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ACCESSION AF237622
VERSION AF237622.1 GI:8164012
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ACCESSION AF327722

VERSION AF327722.1 GI:13195459

KEYWORDS

SOURCE human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 4192)

AUTHORS He, Y.G., Tan, D.Y., Lai, J.H., Xie, Y.F. and Qian, W.

TITLE Cloning and analysis of a human putative acetyltransferase

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 4192)

AUTHORS He, Y.G., Tan, D.Y., Lai, J.H., Xie, Y.F. and Qian, W.

TITLE Direct Submission

JOURNAL Submitted (11-DEC-2000) Biology Department, Yunnan University,
North Street of Greenlake, Kunming, Yunnan 650091, China
FEATURES Location/Qualifiers
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Fluge,O., Bruland,O., Akslen,L.A., Varhaug,J.E. and Lillehaug,J.R.
Identification of NATH, a novel gene overexpressed in papillary
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Unpublished
2 (bases 1 to 5505)
Fluge,O.
Direct Submission
Submitted (31-MAY-2001) Fluge O., Dept. of Molecular Biology,
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1 (bases 1 to 3324)
Choi,S.-C., Kim,J. and Han,J.-K.
Expression of N-terminal acetyltransferase in Xenopus laevis
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2 (bases 1 to 3324)
Choi,S.-C., Kim,J. and Han,J.-K.
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	to NTERMINAL ACETYLTRANSFERRASE 1	(EC 2.3.1.88).			

AK023387
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KEYWORDS: oligo capping; fis (full insert sequence).
SOURCE: Homo sapiens ovary, tumor tissue cDNA to mRNA, clone_lib:OVARC1.
clone:OVARC1001762.

ORGANISM Homo sapiens
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 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (sites)
 AUTHORS Isogai,T., Ota,T., Hayashi,K., Sugiyama,T., Otsuki,T., Suzuki,Y.,
 Nishikawa,T., Nagai,K., Sugano,S., Shiratori,A., Sudo,H.,
 Wagatsuma,M., Hosoiri,T., Kaku,Y., Kodaira,H., Kondo,H.,
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 Ishii,S., Kawai,Y., Saito,K., Yamamoto,J., Wakamatsu,A.,
 Nakamura,Y., Nagahara,K., Masuho,Y., Ninomiya,K. and Iwayanagi,T.
 NEDO human cDNA sequencing project
 NEDO human cDNA sequencing project
 Unpublished (2000)

JOURNAL 2 (bases 1 to 1802)
 REFERENCE Isogai,T. and Otsuki,T.
 AUTHORS Direct Submission
 TITLE Submitted
 JOURNAL Submitted (23-AUG-2000) to the DDBJ/EMBL/GenBank databases. Takao
 Isogai, Helix Research Institute, Genomics Laboratory; 1532-3 Yana,
 Kisarazu, Chiba 292-0812, Japan (E-mail: genomics@hri.co.jp,
 Tel:81-438-52-3951, Fax:81-438-52-3952)

COMMENT NEDO human cDNA sequencing project supported by Ministry of
 International Trade and Industry of Japan; cDNA full insert
 sequencing; Research Association for Biotechnology; cDNA library
 construction, 5'- & 3'-end one pass sequencing and clone selection;;
 Helix Research Institute (supported by Japan Key Technology Center
 etc.) and Department of Virology, Institute of Medical Science,
 University of Tokyo.

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 to N-TERMINAL ACETYLTRANSFERASE 1 (EC 2.3.1.88).
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 VERSION AK023402.1 GI:10435324
 KEYWORDS oligo capping; fis (full insert sequence).
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 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

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VERSION AC015346.1 GI:6435989
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SOURCE fruit fly.
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Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;

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REFERENCE AUTHORS

1 (sites)

Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

Isogai, T., Ota, T., Hayashi, K., Sugiyama, T., Otsuki, T., Suzuki, Y., Nishikawa, T., Nagai, K., Sugano, S., Shiratori, A., Sudo, H., Wagatsuma, M., Takahashi, M., Chiba, Y., Ishida, S., Murakami, K., Ono, Y., Takiguchi, S., Watanabe, S., Kimura, K., Murakami, K., Ishii, S., Kawai, Y., Saito, K., Yamamoto, J., Wakamatsu, A., Nakamura, Y., Nagahara, K., Masuho, Y., Ninomiya, K. and Iwayanagi, T.

NEO human cDNA sequencing project

Unpublished (2000)

2 (bases 1 to 1985)

Isogai, T. and Otsuki, T.

Direct Submission

Submitted (23-AUG-2000) Takao Isogai, Helix Research Institute, Genomics Laboratory, 1532-3 Yana, Kisarazu, Chiba 292-0812, Japan (E-mail: genomics@hri.co.jp. Tel: 81-438-52-3951, Fax: 81-438-52-3952)

NEO human cDNA sequencing project supported by Ministry of International Trade and Industry of Japan; cDNA full insert sequencing; Research Association for Biotechnology; cDNA library construction, 5'- & 3'-end one pass sequencing and clone selection; Helix Research Institute (supported by Japan Key Technology Center etc.) and Department of Virology, Institute of Medical Science, University of Tokyo.

FEATURES

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Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 180213)
AUTHORS Celniker,S.E., Adams,M.D., Kronmiller,B., Tyler,D., Wan,K.H.,
Holt,R.A., Evans,C.A., Gocayne,J.D., Amanatides,P.G., Brandon,R.C.,
Rogers,Y., An,H., Baldwin,D., Banzon,J., Beeson,K.Y., Busam,D.A.,
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McIntosh,T.C., Moy,M., Murphy,B., Nelson,C., Nelson,K.A., Nunoo,J.,
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BACR27L16, complete sequence.

AC011071.12 GI:14249062

HTG.
fruit fly.
Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
pteroygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 182623)
Celniker,S.E., Adams,M.D., Kronmiller,B., Tyler,D., Wan,K.H.,
Holt,R.A., Evans,C.A., Cocayne,J.D., Amanatides,P.G., Brandon,R.C.,
Rogers,Y., An,H., Baldwin,D., Bazzon,J., Beeson,K.Y., Busam,D.A.,
Carlson,J.W., Center,A.L., Champe,M., Davenport,L.B., Dietz,S.M.,
Dodson,K., Dorsett,V., Doyle,C., Dresnek,D., Farfan,D.,
Ferriera,S., Frise,E., Galle,R.F., Garg,N.S., George,R.A.,
Gonzalez,M., Houck,J., Hoskins,R.A., Hostin,D., Howland,T.J.,
Ibegwam,C., Jalali,M., Kruse,D., Li,P., Mattei,B., Moshrefi,A.,
McIntosh,T.C., Moy,M., Murphy,B., Nelson,C., Nelson,K.A., Nunoo,J.,
Pacleb,J., Paragas,V., Park,S., Patel,S., Pfeiffer,B., Scheeler,F.,
Phouanavong,S., Pittman,G.S., Puri,V., Richards,S., Schaefer,F.,
Stapleton,M., Strong,R., Swirskas,R., Tector,C., Williams,S.M.,
Zaveri,J.S., Smith,H.O., Rubin,G.M. and Venter,J.C.
Sequencing of Drosophila chromosome X, region 18C-18D

TITLE
JOURNAL
REFERENCE
AUTHORS

Unpublished
2 (bases 1 to 182623)
Celniker,S.E., Agbayani,A., Arcaluna,T.T., Baxter,E., Blazej,R.G.,
Butenhoff,C., Champe,M., Chavez,C., Chew,M., Ciesiolka,L.,
Doyle,C.M., Farfan,D.E., Galle,R.A., Hummasti,S.R., Karra,K., Kearney,L.,
Hoskins,R.A., Houston,K.A., Lomotan,M.A., Mazda,P.,
Kim,E., Lee,B., Lewis,S., Li,P., Lomotan,M.A., Mazda,P.,
Moshrefi,A.R., Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S.,
Pfeiffer,B., Poen,L., Sequeira,A., Sethi,H., Snir,E., and
Swirskas,R.R., Wan,K.H., Weinburg,T., Zhang,R., Zieran,L.L. and
Rubin,G.M.

TITLE
JOURNAL
COMMENT

Direct Submission
Submitted (01-OCT-1999) Drosophila Genome Center, Lawrence Berkeley
Laboratory, MS 64-121, Berkeley, CA 94720, USA
On May 30, 2001 this sequence version replaced gi:7143400.
Sequence submitted by:
Berkeley Drosophila Genome Project
Lawrence Berkeley National Laboratory, MS 64-121
Berkeley, CA 94720
This sequence was assembled using end sequences from a whole genome
shotgun and from subclones of this BAC and its neighboring clones.
For further information about this sequence, including its location
and relationship to other sequences, please visit our sequence
archive Web site (<http://www.fruitfly.org/sequence/>) or send email
to bdp@fruitfly.berkeley.edu.

FEATURES
source
Location/Qualifiers
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Drosophila melanogaster BAC library, partial EcoRI in
pBACE3.6)"
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BASE COUNT
ORIGIN

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Percent similarity: 69.310 Percent Identity: 42.291

alignment_block:
US-09-836-410A-1 x AC011071

Align seg 1/1 to: AC011071 from: 1 to: 182623

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[illegible]

Mon Jul 22 09:40:54 2002

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